

BEST AVAILABLE COPY

1294933

# THE UNITED STATES OF AMERICA

TO ALL TO WHOM THESE PRESENTS SHALL COME:

UNITED STATES DEPARTMENT OF COMMERCE

United States Patent and Trademark Office

*March 10, 2005*

THIS IS TO CERTIFY THAT ANNEXED HERETO IS A TRUE COPY FROM THE RECORDS OF THE UNITED STATES PATENT AND TRADEMARK OFFICE OF THOSE PAPERS OF THE BELOW IDENTIFIED PATENT APPLICATION THAT MET THE REQUIREMENTS TO BE GRANTED A FILING DATE.

APPLICATION NUMBER: 60/478,185

FILING DATE: *June 13, 2003*

RELATED PCT APPLICATION NUMBER: PCT/US04/19046



Certified by


*Don W. F. Dudas*

Under Secretary of Commerce  
for Intellectual Property  
and Director of the United States  
Patent and Trademark Office

16569 U.S. PTO  
06/13/03

26-160038195-01/1/2003  
SUBSTITUTE PTO/SB/16 (\$-03)

**PROVISIONAL APPLICATION FOR PATENT COVER SHEET**  
This is a request for filing a PROVISIONAL APPLICATION FOR PATENT under 37 CFR §1.53(c).

INVENTOR(S)					
Given Name (first and middle (if any))		Family Name or Surname		Residence (City and either State or Foreign Country)	
Additional inventors are being named on the <u>0</u> separately numbered sheets attached hereto					
TITLE OF THE INVENTION (500 characters max)					
SKN-1 Gene and Protein					
CORRESPONDENCE ADDRESS					
Direct all correspondence to:					
[X] Customer Number: 26161					
					
OR					
[ ] Firm or Individual Name					
Address					
Address					
City					
Country		United States		State	ZIP
				Telephone	Fax
ENCLOSED APPLICATION PARTS (check all that apply)					
[X] Specification	Number of Pages	49	[ ] CD(s), Number		
[X] Drawing(s)	Number of Sheets	31	[ ] Other (specify)		
[ ] Application Data Sheet. See 37 CFR 1.76.					
METHOD OF PAYMENT OF FILING FEES FOR THIS PROVISIONAL APPLICATION FOR PATENT					
[X] Applicant Claims small entity status. See 37 CFR 1.27.					
[X] A check or money order is enclosed to cover the filing fees.					
[ ] The Director is hereby authorized to charge filing fees or credit any overpayment to Deposit Account Number:					
[ ] Payment by credit card. Form PTO-2038 is attached.					
The invention was made by an agency of the United States Government or under a contract with an agency of the United States Government.					
[ ] No.					
[X] Yes, the name of the U.S. Government agency and the Government contract number are:					
National Institutes of Health Grant Nos. RO1GM50900 and RO1GM62891					

Respectfully submitted,

Signature

*John T. Kurall*  
Reg. No. 50,680

Date June 13, 2003

FOR Typed Name John W. Freeman, Esq., Reg. No. 29,066

Telephone No. (617) 542-5070

Docket No. 10861-032P01

20873808.doc

CERTIFICATE OF MAILING BY EXPRESS MAIL

Express Mail Label No. EV331002367

Date of Deposit June 13, 2003

PROVISIONAL APPLICATION FOR PATENT

under

37 CFR §1.53(c)

TITLE: SKN-1 GENE AND PROTEIN

APPLICANT:

CERTIFICATE OF MAILING BY EXPRESS MAIL

Express Mail Label No. EV331002367US

June 13, 2003  
Date of Deposit

## SKN-1 Gene and Protein

### Statement as to Federally Sponsored Research

This invention was made with Government support under National Institutes of Health Grant Nos. RO1GM50900 and RO1GM62891. The Government has certain rights in this invention.

5

### Technical Field

This invention generally relates to the SKN-1 gene and protein, and their use in screening methods for isolating modulators of an oxidative stress response pathway.

### Background

10 In diverse organisms, a common mesendodermal tissue field gives rise to the endoderm and a mesoderm subset that forms the heart and blood in vertebrates. In the nematode *Caenorhabditis elegans*, mesendodermal development is initiated by the maternally expressed transcription factor SKN-1, which specifies the fate of a single cell, the EMS blastomere. The EMS daughter cell E becomes the endoderm, which consists of the intestine. Its sister cell MS gives rise to mesodermal derivatives that include the pharynx, a feeding pump that is analogous  
15 to the heart, and coelomocytes that resemble macrophages. In these embryonic cells SKN-1 induces expression of the GATA factors MED-1 and MED-2, which are required for differentiation of EMS lineages. *C. elegans skn-1* mutants are sensitive to oxidative stress and have shortened lifespans.

20 In vertebrates, Nrf proteins activate transcription of genes encoding the Phase II detoxification enzymes, which constitute the primary cellular defense against oxidative stress. Essentially all organisms must defend themselves against reactive oxygen species (ROS), which are derived from both mitochondrial respiration and exogenous sources. Phase II enzymes synthesize the critical reducing agent glutathione, scavenge ROS directly, and detoxify reactive intermediates that are generated when xenobiotics are metabolized by the cytochrome p450  
25 (Phase I) enzymes. Through Nrf2, exposure to oxidative stress or particular classes of chemicals induces Phase II enzyme gene expression in a variety of tissues, including the liver and digestive tract. This mechanism also constitutes the major response to chemoprotective antioxidants, including many natural compounds, which thereby stimulate xenobiotic detoxification and

inhibit carcinogen-induced tumorigenesis. Accordingly, mice that lack *Nrf2* are abnormally susceptible to drug toxicity and carcinogenesis, and do not respond to chemoprotective antioxidants.

### Summary

5       The present invention is based in part on the discovery that SKN-1 is required for oxidative stress resistance and longevity in *C. elegans*. It has been discovered that SKN-1 orchestrates a major oxidative stress response in *C. elegans*, similar to Nrf proteins in vertebrates, in addition to initiating embryonic development of the *C. elegans* mesendoderm.

10       Accordingly, the present invention provides in vivo methods for determining whether a test compound is a candidate compound capable of modulating the SKN-1-mediated oxidative stress response. For example, in one aspect the invention provides a method for determining whether a test compound is a candidate SKN-1-mediated oxidative stress response-activating compound, comprising: (a) providing a nematode capable of expressing a SKN-1 polypeptide and containing at least one transgene comprising: (i) an oxidative stress resistance gene promoter operably linked to (ii) a reporter gene; and (b) contacting the nematode with the test compound; and (c) determining whether expression of the transgene is increased, wherein an increase in expression of the transgene indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-activating compound.

15       In another aspect, the invention provides a method for determining whether a test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound, comprising: (a) providing a nematode capable of expressing a SKN-1 polypeptide and containing at least one transgene comprising: (i) an oxidative stress resistance gene promoter operably linked to (ii) a reporter gene; (b) contacting the nematode with the test compound; and (c) before, during, or after step (b), subjecting the nematode to conditions that activate the SKN-1-mediated oxidative stress response in the absence of the test compound; and (d) determining whether expression of the transgene is decreased or unchanged, wherein decreased or unchanged expression of the transgene indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound.

20       In still another aspect, the invention provides a method for determining whether a test compound is a candidate SKN-1-mediated oxidative stress response-activating compound,

comprising: (a) providing a nematode containing a transgene encoding a SKN-1 fusion protein, wherein the transgene comprises: (i) a SKN-1 DNA operably linked to (ii) a reporter gene; (b) contacting the nematode with the test compound; and (c) determining whether the SKN-1 fusion protein accumulates in nuclei in the nematode, wherein increased accumulation indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-activating compound.

In still another aspect, the invention provides a method for determining whether a test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound, comprising: (a) providing a nematode containing a transgene encoding a SKN-1 fusion protein, wherein the transgene comprises: (i) a SKN-1 DNA operably linked to (ii) a reporter gene; (b) contacting the nematode with the test compound; (c) before or during step (b), subjecting the nematode to conditions that activate the SKN-1-mediated oxidative stress response in the absence of the test compound; and (d) determining whether the SKN-1 fusion protein accumulates in nuclei in the nematode, wherein decreased or unchanged accumulation of the transgene indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound.

In some embodiments of the inventions described herein, a cultured nematode cell, or cultured mammalian cell is utilized instead of a nematode.

The present invention also provides in vitro methods for determining whether a test compound is a candidate compound capable of modulating the SKN-1-mediated oxidative stress response. In one aspect the invention provides a method for determining whether a test compound is a candidate compound capable of inhibiting a SKN-1-mediated oxidative stress response, comprising: (a) contacting a SKN-1 polypeptide or SKN-1 DNA with a test compound; and (b) detecting interaction of the test compound with the SKN-1 polypeptide SKN-1 DNA, wherein an interaction indicates that the test compound is a candidate compound capable of inhibiting a SKN-1-mediated oxidative stress response.

In another aspect, the invention provides a method for determining whether a test compound is a candidate compound capable of inhibiting a SKN-1-mediated oxidative stress response, comprising: (a) providing a SKN-1 polypeptide or fragment thereof and an oxidative stress resistance gene encoding  $\gamma$ -glutamine cysteine synthase heavy chain, glutathione synthetase, NADH quinone oxidoreductase, superoxide dismutase, catalase, or glutathione S-transferase, or SKN-1 polypeptide-binding fragments thereof; (b) contacting the SKN-1

polypeptide or fragment thereof and the oxidative stress resistance gene or SKN-1 polypeptide-binding fragment thereof with a test compound; and (c) determining whether the SKN-1 polypeptide or fragment thereof and the oxidative stress resistance gene or SKN-1 polypeptide-binding fragment thereof interact in the presence of the test compound, wherein a decrease in  
5 interaction indicates that the test compound is a candidate compound capable of inhibiting a SKN-1-mediated oxidative stress response.

In any of the inventions described herein, the promoter can be a promoter of a gene encoding a protein selected from the group consisting of:  $\gamma$ -glutamine cysteine synthase heavy chain, glutathione synthetase, NADH quinone oxidoreductase, superoxide dismutase, catalase,  
10 and glutathione S-transferase. Further, the reporter gene can be a gene encoding a protein selected from the group consisting of: green fluorescent protein, chloramphenicol acetyl transferase, green fluorescent protein,  $\beta$  glucuronidase, and luciferase. A nematode used in any of the methods described herein can be *Caenorhabditis elegans*.

Further, in embodiments of inventions described herein, a confirmatory step can be  
15 carried out after a candidate compound has been isolated. For example, the confirmatory step can include contacting a nematode (e.g., a second nematode; or a cultured nematode cell, or cultured mammalian cell) with the candidate compound to determine whether the candidate compound decreases or increases oxidative stress resistance, relative to the oxidative stress resistance of the second nematode (or the cultured nematode cell, or the cultured mammalian  
20 cell) not contacted with the candidate compound, wherein a candidate compound that decreases oxidative stress resistance is an oxidative stress response-inhibiting agent, and wherein a candidate compound that increases oxidative stress resistance is an oxidative stress response-activating agent.

In other embodiments, the confirmatory step(s) include: (c) providing a nematode not  
25 capable of expressing a SKN-1 polypeptide (e.g., a *skn-1* mutant) and containing at least one transgene comprising: (i) an oxidative stress resistance gene promoter operably linked to (ii) a reporter gene; and (d) contacting the nematode with the test compound, wherein no increase in expression of the transgene following step (d) indicates that the candidate compound is an oxidative stress response-activating agent.

The present invention also includes compounds capable of activating a SKN-1-mediated oxidative stress response isolated using the methods described herein (e.g., for isolating SKN-1 mediated oxidative stress response inhibitors or activators).

Also within the invention is the use of a compounds identified herein in the manufacture of a medicament for treatment or prevention of a condition described herein. The medicament can be in any form described herein.

Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although suitable methods and materials for the practice or testing of the present invention are described below, other methods and materials similar or equivalent to those described herein, which are well known in the art, can also be used. All publications, patent applications, patents, and other references mentioned herein are incorporated by reference in their entirety. In case of conflict, the present specification, including definitions, will control. In addition, the materials, methods, and examples are illustrative only and not intended to be limiting.

Other features and advantages of the invention will be apparent from the following detailed description, and from the claims.

#### Description of the Drawings

Figs. 1A-1C: SKN-1 embryonic functions and comparison to Nrf proteins. 1A: Cell fate specification. In four-cell embryos, SKN-1 initiates mesendodermal development by establishing the EMS blastomere fate. The anterior pharynx is specified in ABa descendants by a SKN-1-dependent signal from MS. Anterior is to the left, and ventral at the bottom. 1B: SKN-1 compared to Nrf proteins. The SKN-1 minor groove-binding arm is shown in light green. Percent identity between SKN-1 and mouse Nrf2 regions is indicated. 1C: Consensus sequences for SKN-1 binding and the ARE. The SKN-1 BR recognizes a consensus bZIP half-site (underlined) adjacent to an AT-rich motif (grey) that is specified by the arm (B). Nrf proteins bind to the ARE as obligate heterodimers with Maf or other bZIP. R=G/A; W=T/A.



Figs. 2A-2L: *skn-1*-dependent GCS-1::GFP expression in the intestine and ASI neurons. 2A-2F: GCS-1::GFP expression in wild type animals. A *gcs-1* genomic fragment containing its 17 amino terminal codons and 1840 upstream bp was fused to the amino terminus of GFP that contained a nuclear localization signal. Expression patterns shown are each representative of more than two independent transgenic lines, and of all postembryonic stages examined (L2-adult; data not shown). 2A and 2B show Nomarski and fluorescent views, respectively, of an L2 larva. In 2B a line demarcates the approximate boundary between the anterior intestine (I) and posterior pharynx (P). 2C and 2D show combined Nomarski/fluorescent and fluorescent views, respectively, of the head of a typical L4 stage animal that had been exposed to DiI. In 2D one of the two ASI neurons is indicated with an arrow. 2E and 2F show an L2 larva in which GCS-1::GFP expression was induced to high levels in the intestine by heat. A similar induction occurred in response to paraquat (Table 2). The boundary between the anterior intestine and posterior pharynx is indicated as in 2B. 2G-2L: GCS-1::GFP was not detectable outside of the pharynx in *skn-1* homozygotes. Typical animals are shown from experiments which parallel those displayed to the left in 2A-2F. Note the absence of GCS-1::GFP in the intestine and ASI neurons under normal conditions 2G-2J, and after treatment with heat (2K, 2L) or paraquat (data not shown). In two independent transgenic lines, in a homozygous *skn-1* background GCS-1::GFP expression was not detected in these tissues in any animals under either normal or induction conditions.

Figs. 3A-3C: Specific elements required for *skn-1*-independent and -dependent GCS-1::GFP expression. 3A: Analysis of the *gcs-1* promoter. Expression of the indicated constructs from transgenic extrachromosomal arrays was assayed in 2-3 independent transgenic lines, under normal conditions and after induction by paraquat and heat. Approximate relative expression levels in the tissues designated to the right (data not shown) are indicated by + signs, with ++ indicating a reproducible reduction, and + indicating barely detectable expression. Within each set of transgenic lines that carried promoter mutations, levels of normal and induced expression were affected in parallel. Mutations that were created in predicted SKN-1 sites 1, 2, and 3 are described in Materials and Methods, and are not compatible with SKN-1 binding (see text). Red ovals indicate predicted SKN-1 binding sites and a green bar the 5' end of the *gcs-1::gfp* coding region. Map numbers refer to the predicted translation start. 3B: Uncoupling pharyngeal GCS-1::GFP expression from intestinal and ASI neuron expression. The *gcsΔ 2* mutation eliminated

pharyngeal GCS-1::GFP expression, but allowed near-wild type levels of ASI and intestinal expression. Concurrent ablation of SKN-1 binding site 3 (*gcsΔ 2,mut3*) eliminated transgene expression in all tissues. Paraquat-treated worms are shown in the GFP column. 3C: Composite *gcs-1* promoter element that includes SKN-1 site 3, and is also present in the *med-1* and -2 promoters. SKN-1 binding sites are red, and identical sequences are boxed.

Figs. 4A and 4B: Specific binding of SKN-1 to an essential *gcs-1* promoter sequence.

4A: Binding of full-length SKN-1 to site 3 within the *gcs-1* composite element, assayed by EMSA. Lanes 2-5 show binding of increasing amounts of *in vitro* translated SKN-1 protein (0  $\mu$ l, 0.25  $\mu$ l, 0.5  $\mu$ l, 3  $\mu$ l translation lysate; indicated by a triangle) to the wild type site. Lane 1 shows binding to 3  $\mu$ l unprogrammed lysate. A background species is labeled. Lanes 6-10 show the same assay performed with the mutant probe. In lanes 11-20, SKN-1-DNA binding is assayed in the presence of the indicated unlabelled competitor oligonucleotides. Lanes 12-15 and 17-20 correspond to addition of a 20-, 50-, 150-, and 400-fold molar excess of competitor over the labeled wild-type DNA. 4B: The *in vitro* translated SKN-1 DNA binding domain (Fig. 1B) binds specifically to the *gcs-1* composite element. Binding was assayed as in 4A.

Figs. 5A-5G: Expression and stress-induced nuclear accumulation of SKN-1::GFP. 5A: SKN-1::GFP transgenes **a.** *skn-1* gene. Transcribed coding and untranslated regions are indicated in red and blue, respectively. **b.** SKN-1::GFP translational fusion construct, which includes an EcoRI fragment that previously rescued maternal *skn-1* lethality. *C. elegans* DNA is indicated by a black line. **c.** SknPro::GFP promoter fusion, in which the 38 Nterminal SKN-1 amino acids are fused to GFP containing a nuclear localization signal. 5B-5D: Embryonic expression of SKN-1::GFP. 5B, 5C, and 5D show Nomarski (left) and fluorescent (right) views, of 100 cell, 280 min., and three fold embryos, respectively. Endogenous intestinal autofluorescence is visible as yellow or orange. White triangles indicate intestine precursor nuclei. Int: intestine. Ph: pharynx. 5E: SKN-1::GFP expression in ASI neurons (arrows). Nomarski/fluorescent (left) and fluorescent (right) views are shown of a typical DiI-exposed L4 larva. 5F: Larval SKN-1::GFP expression under normal conditions. Fluorescent and Nomarski closeups of the boxed region of this L2 are shown at bottom. Note the low-level SKN-1::GFP expression in intestinal nuclei (white triangle). 5G: SKN-1::GFP localization under oxidative stress. Examination of multiple focal planes revealed that SKN-1::GFP levels were not substantially altered in ASI neurons (arrows), but in many animals were dramatically increased

in intestinal nuclei (Table 3). A heat-shocked L2 is shown, but similar results were obtained upon exposure to other oxidative stress inducers (Table 3). The integrated strain *Is007* is shown, but two extrachromosomal lines and a different integrated line exhibited similar patterns.

Figs. 6A-6B: *skn-1* mutants are sensitive to oxidative stress and have reduced lifespans.

5 6A: Paraquat sensitivity. Worms were scored for survival at the times shown after they had been placed in M9 that contained 100 mM paraquat. An average of three experiments involving 24 worms each is graphed. All wild type and *skn-1* mutant worms survived a parallel control 72 hr. incubation in M9 alone (data not shown). 6B: Lifespan assay. Worms were maintained at 20°C and scored for survival at the indicated time after the L4 stage. An average of three experiments  
10 involving 25-28 worms each is plotted. In wild type, *skn-1(zu67)*, and *skn-1(zu129)* strains, mean life spans were 15.9±2.2, 11.8±1.4, and 11.1±0.1 days, respectively. Mean maximum life spans were 24.3±3.5, 16.3±0.6, and 18.7±0.6 days, respectively.

Fig. 7 illustrates an exemplary regulatory sequence for the glutathione synthetase gene.

Fig. 8 illustrates the sequences of the glutathione synthetase ORF and protein.

15 Fig. 9 illustrates an exemplary regulatory sequence for the NADH quinone oxidoreductase gene.

Fig. 10 illustrates the sequences of the NADH quinone oxidoreductase ORF and protein.

Fig. 11 illustrates an exemplary regulatory sequence for the glutathione S-transferase (R03D7.6) gene.

20 Fig. 12 illustrates the sequences of the glutathione S-transferase (R03D7.6) ORF and protein.

Fig. 13 illustrates an exemplary regulatory sequence for the glutathione S-transferase (F35E8.8) gene.

25 Fig. 14 illustrates the sequences of the glutathione S-transferase (F35E8.8) ORF and protein.

Fig. 15 illustrates an exemplary regulatory sequence for the glutathione S-transferase (F11G11.2) gene.

Fig. 16 illustrates the sequences of the glutathione S-transferase (F11G11.2) ORF and protein.

30 Fig. 17 illustrates an exemplary regulatory sequence for the glutathione S-transferase (K08F4.7) gene.

Fig. 18 illustrates the sequences of the glutathione S-transferase (K08F4.7) ORF and protein.

Fig. 19 illustrates an exemplary regulatory sequence for the superoxide dismutase-1 (*sod-1*) gene.

5 Fig. 20 illustrates the sequences of the superoxide dismutase-1 (*sod-1*) ORF and protein.

Fig. 21 illustrates an exemplary regulatory sequence for the superoxide dismutase-2 (*sod-2*) gene.

Fig. 22 illustrates the sequences of the superoxide dismutase-2 (*sod-2*) ORF and protein.

Fig. 23 illustrates an exemplary regulatory sequence for the catalase (*ctl-1*) gene.

10 Fig. 24 illustrates the sequences of the catalase (*ctl-1*) ORF and protein.

Fig. 25 illustrates an exemplary regulatory sequence for the superoxide dismutase-3 (*sod-3*) gene.

Fig. 26 illustrates the sequences of the superoxide dismutase-3 (*sod-3*) ORF and protein.

15 Fig. 27 illustrates an exemplary regulatory sequence for the  $\gamma$ -glutamine cysteine synthase (also known as glutamate-cysteine ligase) heavy chain gene.

Fig. 28 illustrates the sequences of the  $\gamma$ -glutamine cysteine synthase (also known as glutamate-cysteine ligase) heavy chain open reading frame (ORF) and protein.

Fig. 29 illustrates the sequences of the T19E7.2c SKN-1 ORF and protein.

Fig. 30 illustrates the sequences of the T19E7.2b SKN-1 ORF and protein.

20 Fig. 31 illustrates the sequences of the T19E7.2a SKN-1 ORF and protein.

### DETAILED DESCRIPTION

The present invention relates, in part, to the *C. elegans* SKN-1 gene and protein (a transcription factor), *gcs-1* (encoding  $\gamma$ -glutamine cysteine synthase heavy chain (GCS(h); a target gene thereof) and other oxidative stress resistance genes, e.g., M176.2 (encoding glutathione synthetase); F39B2.3 (encoding NADH quinone oxidoreductase); *sod-1*, *sod-2*, and *sod-3* (encoding superoxide dismutase); *ctl-1* (encoding catalase); and R03D7.6, F35E8.8, F11G11.2, and K08F4.7 (encoding glutathione S-transferase). The invention includes various therapeutic and screening methods. For example, the genes and/or proteins can be used in screening methods to identify, e.g., compounds capable of modulating (e.g., increasing or decreasing) the expression and/or activity of SKN-1, and/or to modulate the oxidative stress

25

30

response pathway in nematodes and other animals (e.g., humans). Such compounds can be used as pharmaceutical agents and/or pesticides.

Before further description of the invention, certain terms employed in the specification, examples, and appended claims are, for convenience, collected here.

5 By "SKN-1-mediated oxidative stress response" is meant an oxidative stress response pathway mediated (i.e., activated) by SKN-1 polypeptides.

The *C. elegans* genome has been sequenced (see, e.g., The *C. elegans* Sequencing Consortium, Science 282, p.2012-2018, 1998), and is accessible through several known electronic databases (see, e.g., the databases accessible at World Wide Web (www) addresses: 10 wormbase.org (WormBase; see, Harris et al., Nucleic Acids Research 31:133-137 (2003), and Stein et al., Nucleic Acids Research 29:82-86 (2001)); ncbi.nlm.nih.gov; and wormbase.sanger.ac.uk). "SKN-1 DNA" or "SKN-1 gene" refers to nucleic acid sequences that include, e.g., the nucleic sequence set forth in Fig. 29 (or the unspliced version thereof)(set forth in the WormBase database as T19E7.2c) and/or Fig. 31 or 30 (or the unspliced versions thereof) 15 (set forth in the WormBase database as T19E7.2a, and T19E7.2b, respectively, homologs thereof, or fragments thereof that encodes SKN-1 polypeptide fragment capable of binding a SKN-1 protein binding site within a promoter of a target gene, e.g., a *C. elegans* Phase II detoxification gene. An example of such a fragment is a fragment that encodes the C-terminal 85 amino acid residues of the SKN-1 polypeptide set forth in Fig. 29 (referred to herein as a "SKN-1 Domain"). By "SKN-1 polypeptide" is meant amino acid sequences that include an amino acid sequence set forth in Figs. 29, 30 and/or 31, or fragments thereof (e.g., the C-terminal 85 amino acid residues of the SKN-1 polypeptide set forth in Fig. 29 (a "SKN-1 Domain)). By "SKN-1 RNA" is meant messenger RNA transcribed from a SKN-1 DNA sequence. 20

25 As used herein, an "oxidative stress resistance gene" is a gene involved in the oxidative stress response in *C. elegans*, and homologs thereof. For example, oxidative stress resistance genes include those demonstrated herein to be targets of the SKN-1 polypeptide, e.g., those encoding  $\gamma$ -glutamine cysteine synthase heavy chain (GCS(h)) (*gcs-1*, set forth in WormBase as F37B12.2); glutathione synthetase (set forth in WormBase as M176.2); NADH quinone oxidoreductase (set forth in WormBase as F39B2.3); superoxide dismutase (*sod-1*, *sod-2*, and *sod-3*; set forth in WormBase as C15F1.7; F10D11.1; C08A9.1, respectively); catalase (*ctl-1*; 30

set forth in WormBase as Y54G11A.6; and several forms of glutathione S-transferase (set forth in WormBase as R03D7.6, F35E8.8, F11G11.2, and K08F4.7), among others.

The sequence of the  $\gamma$ -glutamine cysteine synthase (also known as glutamate-cysteine ligase) heavy chain open reading frame (ORF) and protein are set forth in Fig. 28, and an  
5 exemplary regulatory sequence is set forth in Fig. 27. The sequence of the glutathione synthetase ORF and protein are set forth in Fig. 8, and an exemplary regulatory sequence is set forth in Fig. 7. The sequence of the NADH quinone oxidoreductase ORF and protein are set forth in Fig. 10, and an exemplary regulatory sequence is set forth in Fig. 9. The sequence of one glutathione S-transferase (R03D7.6) ORF and protein are set forth in Fig. 12, and an  
10 exemplary regulatory sequence is set forth in Fig. 11. The sequence of another glutathione S-transferase (F35E8.8) ORF and protein are set forth in Fig. 14, and an exemplary regulatory sequence is set forth in Fig. 13. The sequence of a third glutathione S-transferase (F11G11.2) ORF and protein are set forth in Fig. 16; and an exemplary regulatory sequence is set forth in Fig. 15. The sequence of a fourth glutathione S-transferase (K08F4.7) ORF and protein are set forth in Fig. 18, and an exemplary regulatory sequence is set forth in Fig. 17. The sequence of the superoxide dismutase-1 (*sod-1*) ORF and protein are set forth in Fig. 20, and an exemplary  
15 regulatory sequence is set forth in Fig. 19. The sequence of the superoxide dismutase-2 (*sod-2*) ORF and protein are set forth in Fig. 22 and an exemplary regulatory sequence is set forth in Fig. 21. The sequence of the superoxide dismutase-3 (*sod-3*) ORF and protein are set forth in Fig. 26, and an exemplary regulatory sequence is set forth in Fig. 25. The sequence of the catalase (*ctl-1*) ORF and protein are set forth in Fig. 24 and an exemplary regulatory sequence is set forth in Fig. 23. Predicted SKN-1 binding sites upstream of the genes described above are set forth in Table 1.

A "substantially pure" preparation is a preparation which is at least 60% by weight (dry  
25 weight) the compound of interest, e.g., a SKN-1 polypeptide, or a candidate compound or agent described herein. Preferably the preparation is at least 75%, more preferably at least 90%, and most preferably at least 99%, by weight the compound of interest. Purity can be measured by any appropriate method, e.g., column chromatography, polyacrylamide gel electrophoresis, or HPLC analysis.

30 The term "purified DNA" means DNA that is not immediately contiguous with both of the coding sequences with which it is immediately contiguous (one on the 5' end and one on the

3' end) in the naturally-occurring genome of the organism from which it is derived. The term therefore includes, for example, a recombinant DNA which is incorporated into a vector; into an autonomously replicating plasmid or virus; or into the genomic DNA of a prokaryote or eukaryote, or which exists as a separate molecule (e.g., a cDNA or a genomic DNA fragment produced by PCR or restriction endonuclease treatment) independent of other sequences. It also includes a recombinant DNA that is part of a hybrid gene encoding additional polypeptide sequence.

A "substantially identical" nucleic acid means a nucleic acid sequence which encodes a polypeptide differing only by conservative amino acid substitutions, e.g., substitution of one amino acid for another of the same class (e.g., valine for glycine, arginine for lysine, etc.) or by one or more non-conservative substitutions, deletions, or insertions located at positions of the amino acid sequence which do not destroy the function of the polypeptide.

A "conservative amino acid substitution" is one in which the amino acid residue is replaced with an amino acid residue having a similar side chain. Families of amino acid residues having similar side chains have been defined in the art. These families include amino acids with basic side chains (e.g., lysine, arginine, histidine), acidic side chains (e.g., aspartic acid, glutamic acid), uncharged polar side chains (e.g., glycine, asparagine, glutamine, serine, threonine, tyrosine, cysteine), nonpolar side chains (e.g., alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan), beta-branched side chains (e.g., threonine, valine, isoleucine) and aromatic side chains (e.g., tyrosine, phenylalanine, tryptophan, histidine). Thus, a predicted nonessential amino acid residue can be replaced with another amino acid residue from the same side chain family.

The terms "activate," "induce," "inhibit," "elevate," "increase," "decrease," or the like, denote quantitative differences between two states, e.g., a statistically significant difference, between the two states.

Homology is typically measured using sequence analysis software (e.g., Sequence Analysis Software Package of the Genetics Computer Group, University of Wisconsin Biotechnology Center, 1710 University Avenue, Madison, Wis. 53705, BLAST, or PILEUP/PRETTYBOX programs). Such software matches similar sequences by assigning degrees of homology to various substitutions, deletions, and other modifications.

By "purified antibody" is meant antibody that is at least 60%, by weight, free from the proteins and naturally-occurring organic molecules with which it is naturally associated. Preferably, the preparation is at least 75%, more preferably at least 90%, and most preferably at least 99%, by weight, antibody.

5       The term "longevity" refers to the rate of senescence and/or life-span. The term "immunological methods" refers to any assay involving antibody-based detection techniques including, without limitation, Western blotting, immunoprecipitation, and direct and competitive ELISA and RIA techniques, and "means for detecting" refers to any one or a series of components that sufficiently indicate a detection event of interest. Such means involve at least  
10       one label that may be assayed or observed, including, without limitation, radioactive, fluorescent, and chemiluminescent labels.

By "hybridization techniques" is meant any detection assay involving specific interactions (based on complementarity) between nucleic acid strands, including DNA-DNA, RNA-RNA, and DNA-RNA interactions. Such hybridization techniques may, if desired, include  
15       a PCR amplification step.

The term "modulatory compound", as used herein, refers to any compound capable of affecting (i.e., either increasing or decreasing) SKN-1 expression (i.e., at the level of transcription or translation) or affecting (i.e., either increasing or decreasing) SKN-1 polypeptide activity.

20       The terms "effective amount" and "effective to treat," as used herein, refer to an amount or concentration of a pharmaceutical composition described herein utilized for a period of time (including acute or chronic administration and periodic or continuous administration) that is effective within the context of its administration for causing an intended effect or physiological outcome.

25       The term "patient" is used throughout the specification to describe an animal, human or non-human, to whom treatment according to the methods of the present invention is provided. Veterinary applications are clearly anticipated by the present invention. The term includes but is not limited to birds, reptiles, amphibians, and mammals, e.g., humans, other primates, pigs, rodents such as mice and rats, rabbits, guinea pigs, hamsters, cows, horses, cats, dogs, sheep and  
30       goats. Preferred subjects are humans, farm animals, and domestic pets such as cats and dogs.



The term "treat(ment)," is used herein to denote delaying the onset of, inhibiting, alleviating the effects of, or prolonging the life of a patient.

### Screening Methods

5 In vertebrates, the Nrf pathway has been shown to be critical for oxidative stress resistance, and to be involved in numerous disease states including diabetes, HIV infection, and atherosclerosis, as well as in drug detoxification and cancer chemoprevention. The vertebrate Nrf pathway responds to natural antioxidants found in plants, which produce these compounds for protection against errors in photosynthesis. The present invention is based in part on the  
10 discovery that, in addition to its role in development, SKN-1 plays a role in the *C. elegans* counterpart to the vertebrate Nrf pathway. Like the Nrf pathway, the *C. elegans* pathway mediated by SKN-1 responds to the well-studied natural antioxidant sulforaphane.

The invention provides *in vivo* and *in vitro* screening methods for identifying compounds, e.g., small organic or inorganic molecules (M.W. less than 1,000 Da), oligopeptides,  
15 oligonucleotides, or carbohydrates capable of activating or inhibiting the SKN-1-mediated oxidative stress response.

The screening methods are useful, for example, for isolating novel antioxidants (e.g., compounds that activate the SKN-1-mediated oxidative stress response pathway) or compounds that can be used as pesticides (e.g., compounds that inhibit the SKN-1-mediated oxidative stress  
20 response pathway). Skilled practitioners will also appreciate that the screening methods described herein can be used, for example, to identify or isolate other genes and/or proteins involved in the SKN-1-mediated oxidative stress response pathway, which can themselves be used as pharmaceutical agents or as potential targets for drug discovery.

### In Vivo Screening Methods

25 In one embodiment, the present invention provides a method for determining whether a test compound is a candidate SKN-1-mediated oxidative stress response-activating compound. The method includes providing a nematode, isolated nematode cell, or isolated mammalian cell capable of expressing SKN-1 and containing at least one (i.e., one or more) transgene comprising  
30 an oxidative stress resistance gene promoter operably linked to a reporter gene; and contacting the nematode, isolated nematode cell, or isolated mammalian cell, with the test compound,

wherein an increase in expression of the transgene following contact of the nematode, isolated nematode cell, or isolated mammalian cell, with the test compound indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-activating compound.

5 A similar method can be carried out to determine whether a test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound. The method includes providing a nematode, isolated nematode cell, or isolated mammalian cell capable of expressing SKN-1 and containing at least one (i.e., one or more) transgene comprising an oxidative stress resistance gene promoter operably linked to a reporter gene; contacting the nematode, isolated nematode cell, or isolated mammalian cell, with the test compound; and subjecting the nematode, isolated  
10 nematode cell, or isolated mammalian cell to conditions that activate the SKN-1-mediated oxidative stress response in the absence of the test compound, wherein a decrease or lack of increase in expression of the transgene following the subjecting step indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound.

15 In still another method, a nematode, isolated nematode cell, or isolated mammalian cell capable of expressing a SKN-1 fusion protein is utilized to determine whether a test compound is a candidate SKN-1-mediated oxidative stress response-activating compound. The method includes providing a nematode, isolated nematode cell, or isolated mammalian cell containing a transgene that includes SKN-1 DNA operably linked to a reporter gene; contacting the nematode, isolated nematode cell, or isolated mammalian cell, with the test compound; and determining  
20 whether the SKN-1 fusion protein accumulates in nuclei in the nematode, isolated nematode cell, or isolated mammalian cell. Increased accumulation (e.g., above control levels), indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-activating compound.

25 A similar method can be carried out to determine whether a test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound. The method includes providing a nematode, isolated nematode cell, or isolated mammalian cell containing a transgene that includes SKN-1 DNA operably linked to a reporter gene; contacting the nematode, isolated nematode cell, or isolated mammalian cell, with the test compound; subjecting the nematode, isolated nematode cell, or isolated mammalian cell to conditions that activate the SKN-1-  
30 mediated oxidative stress response in the absence of the test compound; and determining whether the SKN-1 fusion protein accumulates in nuclei in the nematode, isolated nematode cell, or

isolated mammalian cell. Decreased or no increase in accumulation (e.g., as compared to control levels), indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound.

5 A "test compound" can be any chemical compound, for example, a macromolecule (e.g., a polypeptide, a protein complex, or a nucleic acid) or a small molecule (e.g., an amino acid, a nucleotide, an organic or inorganic compound). The test compound can have a formula weight of less than about 10,000 grams per mole, less than 5,000 grams per mole, less than 1,000 grams per mole, or less than about 500 grams per mole. The test compound can be naturally occurring (e.g., an herb or a natural product), synthetic, or can include both natural and synthetic  
10 components. Examples of small molecules include peptides, peptidomimetics (e.g., peptoids), amino acids, amino acid analogs, polynucleotides, polynucleotide analogs, nucleotides, nucleotide analogs, and organic or inorganic compounds, e.g., heteroorganic or organometallic compounds.

15 Test compounds can be screened individually or in parallel. An example of parallel screening is a high throughput drug screen of large libraries of chemicals. Such libraries of test compounds can be generated or purchased, e.g., from Chembridge Corp., San Diego, CA. Libraries can be designed to cover a diverse range of compounds. For example, a library can include 500, 1000, 10,000, 50,000, or 100,000 or more unique compounds. Alternatively, prior experimentation and anecdotal evidence can suggest a class or category of compounds of  
20 enhanced potential. A library can be designed and synthesized to cover such a class of chemicals.

Examples of methods for the synthesis of molecular libraries can be found in the literature, for example in: DeWitt et al. (1993) *Proc. Natl. Acad. Sci. U.S.A.* 90:6909; Erb et al. (1994) *Proc. Natl. Acad. Sci. USA* 91:11422; Zuckermann et al. (1994). *J. Med. Chem.* 37:2678;  
25 Cho et al. (1993) *Science* 261:1303; Carrell et al. (1994) *Angew. Chem. Int. Ed. Engl.* 33:2059; Carell et al. (1994) *Angew. Chem. Int. Ed. Engl.* 33:2061; and Gallop et al. (1994) *J. Med. Chem.* 37:1233.

Libraries of compounds may be presented in solution (e.g., Houghten (1992) *Biotechniques* 13:412-421), or on beads (Lam (1991) *Nature* 354:82-84), chips (Fodor (1993) *Nature* 364:555-556), bacteria (Ladner, U.S. Patent No. 5,223,409), spores (Ladner U.S. Patent  
30 No. 5,223,409), plasmids (Cull et al. (1992) *Proc Natl Acad Sci USA* 89:1865-1869) or on phage

(Scott and Smith (1990) *Science* 249:386-390; Devlin (1990) *Science* 249:404-406; Cwirla *et al.* (1990) *Proc. Natl. Acad. Sci.* 87:6378-6382; Felici (1991) *J. Mol. Biol.* 222:301-310; Ladner *supra.*).

If a nematode is to be used in the present methods, it can be of any genus and species, e.g., any animal, plant, or insect parasitic nematode, or any free-living terrestrial or aquatic (i.e., marine or freshwater) nematode, that is capable of harboring one or more of the constructs described herein. An example of a particularly useful nematode is *C. elegans*. Likewise, if a cultured nematode cell (e.g., a cultured nematode intestinal cell) is to be used, it can be isolated from any genus and species of nematode. Further, if a mammalian cell is to be used, it can be any mammalian cell, e.g., 3T3, HeLa, and/or HD3 erythroblasts, among others.

The oxidative stress resistance gene promoter can be from a gene identified herein to be a target of a SKN-1 polypeptide. As used herein, a "promoter" is a minimal sequence sufficient to direct transcription; the promoter is located in the 5' region of the native gene. For example, the promoter can be from a gene encoding  $\gamma$ -glutamine cysteine synthase heavy chain (GCS(h)) (*gcs-1*, sequences for which are set forth in WormBase as F37B12.2); glutathione synthetase (sequences for which are set forth in WormBase as M176.2); NADH quinone oxidoreductase (set forth in WormBase as F39B2.3); superoxide dismutase (*sod-1*, *sod-2*, and *sod-3*; sequences for which are set forth in WormBase as C15F1.7; F10D11.1; C08A9.1, respectively); catalase (*ctl-1*; sequences for which are set forth in WormBase as Y54G11A.6; and several forms of glutathione S-transferase (sequences for which are set forth in WormBase as R03D7.6, F35E8.8, F11G11.2, and K08F4.7), among others. In certain embodiments of the present invention, a SKN-1 promoter is included in a transgene (sequences for which are set forth in the WormBase database as T19E7.2a, T19E7.2b, and T19E7.2c).

The present invention contemplates that promoters can be modified to provide preferential (i.e., organ- and/or tissue-specific) expression (and/or repression) of a construct. Examples of such modifications are described in detail in Example 1 (below). There, *gcs-1* promoter deletions were constructed using PCR. Predicted SKN-1 polypeptide binding sites (underlined) were altered as follows: Site 1 -608 GATGACAAT to CTGCAGAAT, Site 2 -317 GATGACTTA to CTGCAGTTA, and Site 3 -121 TTTATCATC to TTTCTGCAG. Expression patterns of the construct changed depending upon the deletions made. For example, the *gcsA* 2 mutation eliminated pharyngeal GCS-1::GFP expression, but allowed near-wild type levels of

ASI and intestinal expression. Concurrent ablation of SKN-1 binding site 3 (*gcsA 2,mut3*) eliminated transgene expression in all tissues. Skilled practitioners will appreciate that similar procedures could be used to cause preferential expression of other constructs.

5 The "reporter gene" can be any sequence the expression of which can be detected or measured, other than the coding sequence to which the promoter naturally is operably linked. Typically, the reporter gene is heterologous to the nematode, isolated nematode cell, or isolated mammalian cell in which promoter activity is measured. Examples of reporter genes include, without limitation, genes that encode green fluorescent protein (or any other fluorescent marker), chloramphenicol acetyl transferase (*cat*),  $\beta$ -glucuronidase (*gus*),  $\beta$ -Galactosidase (*lacZ*),  
10 luciferase, and the like. Reporter gene expression can be measured by any of a number of conventional methods, and the optimal method will depend upon factors such as the nature and function of the reporter gene. In general, suitable assays of reporter gene expression include methods such as (i) assaying the function of a product of the reporter gene (e.g., measuring an enzymatic reaction catalyzed by a product of the reporter gene); (ii) measuring the level of  
15 protein expressed from the reporter gene (e.g., by SDS-PAGE or in an immunoassay using antibodies (e.g., polyclonal or monoclonal antibodies) that specifically bind to the product of the reporter gene; and (iii) measuring the level of mRNA transcribed from the reporter gene. Included within the invention are assays that permit high throughput screening of test compounds.

20 The aforementioned reporter genes, and methods for measuring their expression, are well known to those of ordinary skill in the art. Methods for making the transgenes and their incorporation into the genomes of organisms, e.g., nematodes, or cells, e.g., cultured nematode or mammalian cells, are also well known in the art. Example 1 (below) describes in detail how to make *gcs-1::gfp* and *skn-1::gfp* transgenes, how to incorporate such transgenes into the genome  
25 of *C. elegans*, and how to measure expression of the transgenes (a microscopy-based screen is performed in that Example). Skilled practitioners will appreciate that similar protocols could be used to create other transgenes described herein, and to incorporate those transgenes into other types of nematodes and cultured cells. The present invention contemplates that more than one type of transgene can be inserted into the genome of a nematode or cell.

30 Compounds capable of activating the SKN-1-mediated oxidative stress response in an initial screen discussed above can be considered candidate activating compounds, i.e., candidate

SKN-1-mediated oxidative stress response-activating compounds. Such candidate compounds can be subjected to a confirmatory step, e.g., to determine whether the candidate compound increases the overall oxidative stress resistance of a nematode or cultured cell (e.g., a mammalian cell). Methods for evaluating the oxidative stress resistance of a cell or organism (e.g., relative to controls) are well known in the art, and include, for example, subjecting an organism or cell to conditions of oxidative stress (e.g., using known compounds or by increasing the temperature of the culture environment) and measuring overall survival rate. Alternatively or in addition, the confirmatory step can involve determining whether the candidate compound is capable of increasing expression of at least one gene involved in the vertebrate Nrf oxidative stress response pathway, using transgenes (similar to those described herein) or any other method known in the art for measuring increased expression (e.g., Western blotting). Candidate activating compounds that increase oxidative stress resistance in a nematode, cultured nematode cell, or mammalian cell, or that are capable of increasing expression of at least one gene involved in the vertebrate Nrf oxidative stress response pathway, can be considered oxidative stress response-activating agents.

Alternatively or in addition, a different confirmatory step can be carried out. This step involves providing a nematode not capable of expressing a SKN-1 polypeptide (i.e., a *skn-1* mutant) containing at least one transgene that includes an oxidative stress resistance gene promoter operably linked to a reporter gene; and contacting the nematode with the test compound. If the transgene displays no increase in expression, the candidate compound specifically activates the SKN-1-mediated oxidative stress response, and can be considered an oxidative stress response-activating agent.

Compounds capable of inhibiting the SKN-1-mediated oxidative stress response in an initial screen discussed above can be considered candidate inhibiting compounds, i.e., candidate SKN-1-mediated oxidative stress response-inhibiting compounds. Such candidate inhibitory compounds can also be subjected to a confirmatory step, e.g., to determine whether the candidate inhibitory compound decreases the overall oxidative stress resistance of a nematode or cultured cell (e.g., a mammalian cell). Alternatively or in addition, the confirmatory step can involve determining whether the candidate compound is capable of decreasing (or preventing increased expression under conditions of oxidative stress) expression of at least one gene involved in the vertebrate Nrf oxidative stress response pathway (or an oxidative stress resistance gene not

utilized in the initial screen). Such screens can be carried out using transgenes similar to those described herein or any other method known in the art for measuring increased or decreased expression (e.g., Western blotting). Candidate inhibiting compounds that decrease oxidative stress resistance in a nematode, cultured nematode cell, or mammalian cell, or prevent (or  
5 decrease) increased expression of another oxidative stress resistance gene under conditions of oxidative stress, can be considered an oxidative stress response-inhibiting agent.

Some screens of the present invention require subjecting the nematode or cell to conditions that activate the SKN-1-mediated oxidative stress response in the absence of a test compound. Such conditions include exposing the nematode or cell to a known antioxidant, e.g.,  
10 sulforaphane. Other conditions include, e.g., exposure to the herbicide paraquat (methyl viologen), heat, CdCl<sub>2</sub>, arsenite, H<sub>2</sub>O<sub>2</sub>, diamide, and/or sodium azide.

#### In Vitro Screening Methods

The invention also provides *in vitro* methods for determining whether test compound is a  
15 candidate SKN-1-mediated oxidative stress response-inhibiting compound. In one embodiment, the method includes contacting a SKN-1 polypeptide, or a fragment thereof (e.g., a SKN-1 Domain) or SKN-1 DNA with a test compound; and detecting interaction of the test compound with the SKN-1 polypeptide (or fragment thereof) or SKN-1 DNA. An interaction (e.g., direct or indirect binding) indicates that the test compound is a candidate SKN-1-mediated oxidative  
20 stress response-inhibiting compound.

Another method takes advantage of interactions between SKN-1 polypeptides and oxidative stress resistance genes that are targets of SKN-1 polypeptides. The method includes providing a polypeptide that includes a SKN-1 polypeptide (or fragment thereof that is capable of interacting (i.e., binding) with an oxidative stress resistance gene, e.g., a SKN-1 Domain), and  
25 an oxidative stress resistance gene. The oxidative stress resistance gene can be a gene identified herein as being a target for SKN-1 polypeptides, e.g., genes encoding  $\gamma$ -glutamine cysteine synthase heavy chain, glutathione synthetase, NADH quinone oxidoreductase, superoxide dismutase, catalase, or glutathione S-transferase. Alternatively, a nucleic acid sequence that includes a SKN-1 polypeptide-binding fragment of the oxidative stress resistance gene can be  
30 provided. Examples of SKN-1 polypeptide binding fragments for a number of oxidative stress resistance genes are provided in Table 1. The method further includes contacting the polypeptide comprising the SKN-1 polypeptide (or fragment thereof) and a nucleotide sequence comprising

an oxidative stress resistance gene (or SKN-1 polypeptide-binding fragment thereof) with a test compound; and determining whether the SKN-1 polypeptide (or fragment thereof) and the oxidative stress resistance gene (or SKN-1 polypeptide-binding fragment thereof) interact (i.e., bind) in the presence of the test compound. If no or decreased interaction is evident, the test compound can be considered a candidate SKN-1-mediated oxidative stress response-inhibiting compound.

As in the in vivo screening methods, compounds isolated using the in vitro methods can be subjected to any confirmatory step herein described herein. Skilled practitioners will appreciate that in any screening method described herein, homologues of SKN-1 (e.g., genes or polypeptides) could be substituted for the SKN-1 DNA or polypeptide. Where such substitutions are made, the screens could be carried out essentially as described herein.

#### Medicinal Chemistry

Once a compound (or agent) of interest has been identified, standard principles of medicinal chemistry can be used to produce derivatives of the compound. Derivatives can be screened for improved pharmacological properties, for example, efficacy, pharmaco-kinetics, stability, solubility, and clearance. The moieties responsible for a compound's activity in the assays described above can be delineated by examination of structure-activity relationships (SAR) as is commonly practiced in the art. A person of ordinary skill in pharmaceutical chemistry could modify moieties on a lead compound and measure the effects of the modification on the efficacy of the compound to thereby produce derivatives with increased potency. For an example, see Nagarajan *et al.* (1988) *J. Antibiot.* 41: 1430-8. Furthermore, if the biochemical target of the compound (or agent) is known or determined, the structure of the target and the compound can inform the design and optimization of derivatives. Molecular modeling software is commercially available (e.g., Molecular Simulations, Inc.) for this purpose.

#### Isolated Compounds

Skilled practitioners will appreciate that compounds isolated using the screens described herein can be used to treat a number of conditions or disorders.

Oxidative stress contributes to human pathologies that include diabetes, atherosclerosis, neurodegenerative diseases, reperfusion injury, and HIV infection (see, e.g., Finkel *et al.*, *Nature* 408: 239-247 (2000)). The ROS defenses mobilized by human Nrf proteins are thought to be



beneficial in these diverse disease states. This gene activation pathway is also important for drug detoxification, and therefore for chemotherapeutic agent tolerance, and it may provide a widely applicable means of cancer prevention (see, e.g., Chan et al. *Proc Natl Acad Sci U.S.A* **98**: 4611-6. (2001); Hayes et al., *Cancer Lett* **174**: 103-113 (2001); and Wolf *Proc Natl Acad Sci U.S.A* **98**: 2941-2943 (2001)). For example, dietary consumption of chemoprotective antioxidants acts through Nrf2 to inhibit chemical carcinogenesis in mice, and decreases the risk of gastrointestinal and lung tumors in humans (see, e.g., Ramos-Gomez et al. *Proc Natl Acad Sci U.S.A* **98**: 3410-3415 (2001); Fahey et al. *Proc Natl Acad Sci U.S.A* **99**: 7610-7615 (2002); and Thimmulappa et al. *Cancer Res* **62**: 5196-5203 (2002)).

Accordingly, candidate SKN-1-mediated oxidative stress response-activating compounds shown to increase the oxidative stress resistance of mammalian cells (oxidative stress response-activating agents) can be used as novel antioxidants. Such antioxidants could be used to treat a number of conditions including, but not limited to, aging, cancer (e.g., wherein such an agent can be used in chemoprevention), arteriosclerosis, the effects of diabetes (e.g., the neuropathy and vascular complications associated therewith, islet cell destruction, and detrimental insulin responses), neurodegenerative diseases (e.g., by increasing neuronal oxidative stress resistance and, therefore, survival), reperfusion injury (e.g., injury arising from oxidative stress caused by hypotension, myocardial infarction, and/or stroke); the effects of sleep apnea (e.g., vascular injury arising from the cycle of hypoxia/reoxygenation); viral infection (e.g., human immunodeficiency virus infection); bacterial infections (e.g., in the gut); and toxicity (e.g., drug toxicity (e.g., arising from chemotherapy), heavy metal toxicity, and hepatic toxicity).

Inhibitors are also useful in certain clinical settings. For example, candidate SKN-1-mediated oxidative stress response-inhibiting compounds shown to reduce the oxidative stress resistance of mammalian cells (oxidative stress response-inhibiting agents) can be used to eliminate undesirable cells in an animal, e.g., tumor cells in a human. Further, in certain cancer cells, the oxidative stress response pathway is up-regulated, conferring growth advantages. An inhibitor could be used as a component of a chemotherapeutic regimen to prevent such up-regulation. As still another example, reactive oxygen species (ROS) production plays a detrimental role in certain inflammatory responses, which could be prevented or decreased using an inhibitor described herein.

### Modulation of the SKN-1-Mediated Oxidative Stress Response

The invention provides methods for modulating the SKN-1-mediated oxidative stress response that fall into two basic categories: inhibiting (e.g., limiting or reducing) the SKN-1 SKN-1-mediated oxidative stress response, and activating, e.g., supplementing or providing oxidative stress response activity where there is insufficient or no activity. Whether the SKN-1 mediated oxidative stress response is inhibited or increased depends on the intended application.

#### Inhibition of the Response

In some embodiments, the invention provides for inhibiting the SKN-1-mediated oxidative stress response. Agents that inhibit can be used as, e.g., novel pesticides to control insects or nematodes (e.g., pathogenic nematodes). Agents that inhibit the SKN-1-mediated oxidative stress response are useful to inhibit nematode reproduction, decrease nematode lifespan, and increase nematode sensitivity to oxidative stresses (thereby making the nematode easier to eradicate, e.g., using known pesticides). Such agents are also useful for inhibiting SKN-1 activation of the oxidative stress response in a laboratory/research setting in order to identify other genes and/or proteins involved in this oxidative stress response pathway.

For example, an antisense nucleic acid effective to inhibit expression of an endogenous SKN-1 gene can be utilized. The antisense nucleic acid can include a nucleotide sequence complementary to an entire SKN-1 RNA or only a portion of the RNA. On one hand, the antisense nucleic acid needs to be long enough to hybridize effectively with the SKN-1 RNA. Therefore, the minimum length is approximately 10, 11, 12, 13, 14, or 15 nucleotides. On the other hand, as length increases beyond about 150 nucleotides, effectiveness at inhibiting translation increases only marginally, while difficulty in introducing the antisense nucleic acid into target cells may increase significantly. In view of these considerations, a preferred length for the antisense nucleic acid is from about 15 to about 150 nucleotides, e.g., 20, 25, 30, 35, 40, 45, 50, 60, 70, or 80 nucleotides. The antisense nucleic acid can be complementary to a coding region of SKN-1 mRNA or a 5' or 3' non-coding region of a SKN-1 mRNA (or both). One approach is to design the antisense nucleic acid to be complementary to a region on both sides of the translation start site of the SKN-1 mRNA.

The antisense nucleic acid can be chemically synthesized, e.g., using a commercial nucleic acid synthesizer according to the vendor's instructions. Alternatively, the antisense

nucleic acids can be produced using recombinant DNA techniques. An antisense nucleic acid can incorporate only naturally occurring nucleotides. Alternatively, it can incorporate variously modified nucleotides or nucleotide analogs to increase its *in vivo* half-life or to increase the stability of the duplex formed between the antisense molecule and its target RNA. Examples of nucleotide analogs include phosphorothioate derivatives and acridine-substituted nucleotides. Given the description of the targets and sequences, the design and production of suitable antisense molecules is within ordinary skill in the art. For guidance concerning antisense nucleic acids, see, e.g., Goodchild, "Inhibition of Gene Expression by Oligonucleotides," in *Topics in Molecular and Structural Biology, Vol. 12: Oligodeoxynucleotides* (Cohen, ed.), MacMillan Press, London, pp. 53-77.

Delivery of antisense oligonucleotides can be accomplished by any method known to those of skill in the art. For example, delivery of antisense oligonucleotides for cell culture and/or *ex vivo* work can be performed by standard methods such as the liposome method or simply by addition of membrane-permeable oligonucleotides. To resist nuclease degradation, chemical modifications such as phosphorothionate backbones can be incorporated into the molecule.

Delivery of antisense oligonucleotides for *in vivo* applications can be accomplished, for example, via local injection of the antisense oligonucleotides at a selected site. This method has previously been demonstrated for psoriasis growth inhibition and for cytomegalovirus inhibition. See, for example, Wraight et al., (2001). *Pharmacol Ther.* Apr; 90(1):89-104.; Anderson, et al., (1996) *Antimicrob Agents Chemother* 40: 2004-2011; and Crooke et al., *J Pharmacol Exp Ther* 277: 923-937.

Similarly, the present invention anticipates that RNA interference (RNAi) techniques could be used to inhibit the SKN-1-mediated oxidative stress response, in addition or as an alternative to, the use of antisense techniques. For example, small interfering RNA (siRNA) duplexes directed against SKN-1, or any oxidative stress response gene target of SKN-1, could be synthesized and used to prevent expression of the encoded protein(s). Skilled practitioners will also appreciate that RNAi techniques could be used in screens to identify other genes and/or proteins that modulate the SKN-1 oxidative stress response pathway. For example, these techniques could be used in a screen for genes that when inhibited allowed constitutive activation

of the *gcs-1::gfp* transgene, or that prevented the *gcs-1::gfp* transgene from being activated by oxidative stress or antioxidants.

As another example, SKN-1 polypeptide activity can be inhibited using a SKN-1 polypeptide binding molecule such as an antibody, e.g., an anti-SKN-1 polypeptide antibody, or a SKN-1 polypeptide-binding fragment thereof. The anti-SKN-1 polypeptide antibody can be a polyclonal or a monoclonal antibody. Alternatively or in addition, the antibody can be produced recombinantly, e.g., produced by phage display or by combinatorial methods as described in, e.g., Ladner et al. U.S. Patent No. 5,223,409; Kang et al. International Publication No. WO 92/18619; Dower et al. International Publication No. WO 91/17271; Winter et al. International Publication WO 92/20791; Markland et al. International Publication No. WO 92/15679; Breitling et al. International Publication WO 93/01288; McCafferty et al. International Publication No. WO 92/01047; Garrard et al. International Publication No. WO 92/09690; Ladner et al. International Publication No. WO 90/02809; Fuchs et al. (1991) *Bio/Technology* 9:1370-1372; Hay et al. (1992) *Hum Antibod Hybridomas* 3:81-85; Huse et al. (1989) *Science* 246:1275-1281; Griffiths et al. (1993) *EMBO J* 12:725-734; Hawkins et al. (1992) *J Mol Biol* 226:889-896; Clackson et al. (1991) *Nature* 352:624-628; Gram et al. (1992) *PNAS* 89:3576-3580; Garrad et al. (1991) *Bio/Technology* 9:1373-1377; Hoogenboom et al. (1991) *Nuc Acid Res* 19:4133-4137; and Barbas et al. (1991) *PNAS* 88:7978-7982.

As used herein, the term "antibody" refers to a protein comprising at least one, and preferably two, heavy (H) chain variable regions (abbreviated herein as VH), and at least one and preferably two light (L) chain variable regions (abbreviated herein as VL). The VH and VL regions can be further subdivided into regions of hypervariability, termed "complementarity determining regions" ("CDR"), interspersed with regions that are more conserved, termed "framework regions" (FR). The extent of the framework region and CDR's has been precisely defined (see, Kabat, E.A., et al. (1991) *Sequences of Proteins of Immunological Interest, Fifth Edition*, U.S. Department of Health and Human Services, NIH Publication No. 91-3242, and Chothia, C. et al. (1987) *J. Mol. Biol.* 196:901-917). Each VH and VL is composed of three CDR's and four FRs, arranged from amino-terminus to carboxy-terminus in the following order: FR1, CDR1, FR2, CDR2, FR3, CDR3, FR4.

An anti-SKN-1 polypeptide antibody can further include a heavy and light chain constant region, to thereby form a heavy and light immunoglobulin chain, respectively. The antibody can

be a tetramer of two heavy immunoglobulin chains and two light immunoglobulin chains, wherein the heavy and light immunoglobulin chains are inter-connected by, e.g., disulfide bonds. The heavy chain constant region is comprised of three domains, CH1, CH2, and CH3. The light chain constant region is comprised of one domain, CL. The variable region of the heavy and light chains contains a binding domain that interacts with an antigen. The constant regions of the antibodies typically mediate the binding of the antibody to host tissues or factors, including various cells of the immune system (e.g., effector cells) and the first component (C1q) of the classical complement system.

A "SKN-1 polypeptide-binding fragment" of an antibody refers to one or more fragments of a full-length antibody that retain the ability to specifically bind to SKN-1 polypeptide or a portion thereof. "Specifically binds" means that an antibody or ligand binds to a particular target to the substantial exclusion of other substances. Examples of SKN-1 polypeptide binding fragments of an anti-SKN-1 polypeptide antibody include, but are not limited to: (i) a Fab fragment, a monovalent fragment consisting of the VL, VH, CL and CH1 domains; (ii) a F(ab')<sub>2</sub> fragment, a bivalent fragment comprising two Fab fragments linked by a disulfide bridge at the hinge region; (iii) a Fd fragment consisting of the VH and CH1 domains; (iv) a Fv fragment consisting of the VL and VH domains of a single arm of an antibody, (v) a dAb fragment (Ward *et al.*, (1989) *Nature* 341:544-546), which consists of a VH domain; and (vi) an isolated complementarity determining region (CDR). Furthermore, although the two domains of the Fv fragment, VL and VH, are encoded by separate genes, they can be joined, using recombinant methods, by a synthetic linker that enables them to be made as a single protein chain in which the VL and VH regions pair to form monovalent molecules (known as single chain Fv (scFv); see *e.g.*, Bird *et al.* (1988) *Science* 242:423-426; and Huston *et al.* (1988) *Proc. Natl. Acad. Sci. USA* 85:5879-5883). Such single chain antibodies are also encompassed within the term "SKN-1 polypeptide-binding fragment" of an antibody. These antibody fragments can be obtained using conventional techniques known to those with skill in the art.

The anti-SKN-1 polypeptide antibody can be a fully human antibody (e.g., an antibody made in a mouse which has been genetically engineered to produce an antibody from a human immunoglobulin sequence), or a non-human antibody, e.g., a rodent (mouse or rat), goat, primate (e.g., monkey), camel, donkey, porcine, or fowl antibody.

An anti-SKN-1 polypeptide antibody can be one in which the variable region, or a portion thereof, e.g., the CDRs, are generated in a non-human organism, e.g., a rat or mouse. The anti-SKN-1 polypeptide antibody can also be, for example, chimeric, CDR-grafted, or humanized antibodies. The anti-SKN-1 polypeptide antibody can also be generated in a non-  
5 human organism, e.g., a rat or mouse, and then modified, e.g., in the variable framework or constant region, to decrease antigenicity in a human.

Another approach to inhibiting SKN-1 activity is the administration of a SKN-1 antagonist that binds to (i.e., blocks) SKN-1 polypeptides and prevents it from interacting with its target gene (e.g., a gene involved in the oxidative stress response, e.g., a Phase II  
10 detoxification gene). Such SKN-1 polypeptide antagonists can be identified using a screening method described herein. Alternatively, the SKN-1 antagonist can be an anti-SKN-1 polypeptide antibody, or fragment thereof, as described above.

#### Activation of the Response

15 In some embodiments, the invention provides for activating the SKN-1-mediated oxidative stress response. Agents that activate can be used, e.g., to increase the oxidative stress resistance of beneficial pathogenic nematodes, e.g., those used to protect crops by eliminating/controlling the population of certain insects. Examples of such nematodes are  
20 *Steinernema carpocapsae*, *S. Glaseri* and *Heterorhabditis* spp. Such agents are also useful for, e.g., activating the oxidative stress response via SKN-1 in a laboratory/research setting in order to identify other genes and/or proteins involved in the oxidative stress response pathway.

For example, new or supplemental SKN-1 activity can be provided in vivo by direct administration of a recombinant SKN-1 polypeptide, e.g., to pathogenic nematodes prior to, during, and/or after their introduction into the environment of interest. SKN-1 polypeptides that  
25 can be used to supplemental SKN-1 activity are described herein, e.g., SEQ ID NO:2, or a fragment thereof. Another is described in Example 1. There, a SKN-1/green fluorescent protein fusion protein is described. Such polypeptides can be used "as is" or modified. Examples of modifications include derivation of amino acid side chains, glycosylation, conservative amino acid substitutions, and chemical conjugation or fusion to other non-SKN-1 polypeptide moieties.

30 Alternatively or in addition, a SKN-1 polypeptide can be introduced indirectly into an organism, e.g., a nematode, by expressing within the cells of the organism a nucleic acid

construct containing a nucleotide sequence encoding a SKN-1 polypeptide. Any appropriate expression vector suitable for transfecting the cells of the organism of interest can be used in the invention. The nucleic acid construct can be derived from a non-replicating linear or circular DNA or RNA vector, or from an autonomously replicating plasmid or viral vector. Methods for constructing suitable expression vectors are known in the art, and useful materials are commercially available. With respect to nematodes, an example for producing transgenic *skn-1::gfp* strains of *C. elegans* is discussed in detail in Example 1.

Another approach to increasing SKN-1 activity is the administration of an antioxidant (e.g., sulforaphane) or other compound, e.g., a compound isolated using one of the screening methods described above. Such a compound can be, e.g., a small organic or inorganic molecule, e.g., a novel antioxidant.

### Pharmaceutical Compositions

The compounds, nucleic acids, and polypeptides, fragments thereof, as well as antibodies, e.g., anti-SKN-1 polypeptide antibodies other molecules and agents of the invention (also referred to herein as "active compounds"), e.g., novel antioxidants, can be incorporated into pharmaceutical compositions. Such compositions typically include the nucleic acid molecule, protein, or antibody and a pharmaceutically acceptable carrier. A "pharmaceutically acceptable carrier" can include solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like, compatible with pharmaceutical administration. Supplementary active compounds can also be incorporated into the compositions.

A pharmaceutical composition is formulated to be compatible with its intended route of administration. Examples of routes of administration include parenteral, e.g., intravenous, intradermal, subcutaneous, oral (e.g., inhalation), transdermal (topical), transmucosal, and rectal administration. Solutions or suspensions used for parenteral, intradermal, or subcutaneous application can include the following components: a sterile diluent such as water for injection, saline solution, fixed oils, polyethylene glycols, glycerine, propylene glycol or other synthetic solvents; antibacterial agents such as benzyl alcohol or methyl parabens; antioxidants such as ascorbic acid or sodium bisulfite; chelating agents such as ethylenediaminetetraacetic acid; buffers such as acetates, citrates or phosphates and agents for the adjustment of tonicity such as sodium chloride or dextrose. pH can be adjusted with acids or bases, such as hydrochloric acid

or sodium hydroxide. The parenteral preparation can be enclosed in ampoules, disposable syringes or multiple dose vials made of glass or plastic.

Pharmaceutical compositions suitable for injectable use include sterile aqueous solutions (where water soluble) or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersion. For intravenous administration, suitable carriers include physiological saline, bacteriostatic water, Cremophor EL™ (BASF, Parsippany, NJ) or phosphate buffered saline (PBS). In all cases, the composition must be sterile and should be fluid to the extent that easy syringability exists. It should be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. Prevention of the action of microorganisms can be achieved by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, ascorbic acid, thimerosal, and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars, polyalcohols such as mannitol, sorbitol, sodium chloride in the composition. Prolonged absorption of the injectable compositions can be achieved by including an agent which delays absorption, e.g., aluminum monostearate and gelatin in the composition.

Sterile injectable solutions can be prepared by incorporating the active compound in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the active compound into a sterile vehicle which contains a basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and freeze-drying which yields a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof.

Oral compositions generally include an inert diluent or an edible carrier. For the purpose of oral therapeutic administration, the active compound can be incorporated with excipients and used in the form of tablets, troches, or capsules, e.g., gelatin capsules. Oral compositions can



also be prepared using a fluid carrier for use as a mouthwash. Pharmaceutically compatible binding agents, and/or adjuvant materials can be included as part of the composition. The tablets, pills, capsules, troches and the like can contain any of the following ingredients, or compounds of a similar nature: a binder such as microcrystalline cellulose, gum tragacanth or gelatin; an excipient such as starch or lactose, a disintegrating agent such as alginic acid, Primogel, or corn starch; a lubricant such as magnesium stearate or Sterotes; a glidant such as colloidal silicon dioxide; a sweetening agent such as sucrose or saccharin; or a flavoring agent such as peppermint, methyl salicylate, or orange flavoring.

For administration by inhalation, the compounds are delivered in the form of an aerosol spray from pressured container or dispenser that contains a suitable propellant, e.g., a gas such as carbon dioxide, or a nebulizer.

Systemic administration can also be by transmucosal or transdermal means. For transmucosal or transdermal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art, and include, for example, for transmucosal administration, detergents, bile salts, and fusidic acid derivatives. Transmucosal administration can be accomplished through the use of nasal sprays or suppositories. For transdermal administration, the active compounds are formulated into ointments, salves, gels, or creams as generally known in the art.

The compounds can also be prepared in the form of suppositories (e.g., with conventional suppository bases such as cocoa butter and other glycerides) or retention enemas for rectal delivery.

In one embodiment, the active compounds are prepared with carriers that will protect the compound against rapid elimination from the body, such as a controlled release formulation, including implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Methods for preparation of such formulations will be apparent to those skilled in the art. The materials can also be obtained commercially from Alza Corporation and Nova Pharmaceuticals, Inc. Liposomal suspensions (including liposomes targeted to infected cells with monoclonal antibodies to viral antigens) can also be used as pharmaceutically acceptable carriers. These can be prepared according to methods known to those skilled in the art, for example, as described in U.S. Patent No. 4,522,811.

It is advantageous to formulate oral or parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the subject to be treated; each unit containing a predetermined quantity of active compound calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier.

Toxicity and therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, *e.g.*, for determining the LD50 (the dose lethal to 50% of the population) and the ED50 (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD50/ED50. Compounds which exhibit high therapeutic indices are preferred. While compounds that exhibit toxic side effects may be used, care should be taken to design a delivery system that targets such compounds to the site of affected tissue, *e.g.*, bone or cartilage, in order to minimize potential damage to uninfected cells and, thereby, reduce side effects.

The data obtained from cell culture assays and animal studies can be used in formulating a range of dosage for use in humans. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED50 with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. For any compound used in the method of the invention, the therapeutically effective dose can be estimated initially from cell culture assays. A dose may be formulated in animal models to achieve a circulating plasma concentration range that includes the IC50 (*i.e.*, the concentration of the test compound which achieves a half-maximal inhibition of symptoms) as determined in cell culture. Such information can be used to more accurately determine useful doses in humans. Levels in plasma may be measured, for example, by high performance liquid chromatography.

For the SKN-1 modulating agents described herein, an effective amount, *e.g.* of a protein or polypeptide (*i.e.*, an effective dosage), ranges from about 0.001 to 30 mg/kg body weight, *e.g.* about 0.01 to 25 mg/kg body weight, *e.g.* about 0.1 to 20 mg/kg body weight. The protein or polypeptide can be administered one time per week for between about 1 to 10 weeks, *e.g.* between 2 to 8 weeks, about 3 to 7 weeks, or for about 4, 5, or 6 weeks. The skilled artisan will appreciate that certain factors influence the dosage and timing required to effectively treat a

patient, including but not limited to the type of patient to be treated, the severity of the disease or disorder, previous treatments, the general health and/or age of the patient, and other diseases present. Moreover, treatment of a patient with a therapeutically effective amount of a protein, polypeptide, antibody, or other compound can include a single treatment or, preferably, can  
5 include a series of treatments.

For antibodies, a useful dosage is 0.1 mg/kg of body weight (generally 10 mg/kg to 20 mg/kg). Generally, partially human antibodies and fully human antibodies have a longer half-life within the human body than other antibodies. Accordingly, lower dosages and less frequent administration are possible. Modifications such as lipidation can be used to stabilize antibodies  
10 and to enhance uptake and tissue penetration. A method for lipidation of antibodies is described by Cruikshank *et al.* ((1997) *J. Acquired Immune Deficiency Syndromes and Human Retrovirology* 14:193).

If the agent is a small molecule, exemplary doses include milligram or microgram amounts of the small molecule per kilogram of subject or sample weight (e.g., about 1  
15 microgram per kilogram to about 500 milligrams per kilogram, about 100 micrograms per kilogram to about 5 milligrams per kilogram, or about 1 microgram per kilogram to about 50 micrograms per kilogram. It is furthermore understood that appropriate doses of a small molecule depend upon the potency of the small molecule with respect to the expression or activity to be modulated. When one or more of these small molecules is to be administered to an  
20 animal (e.g., a human) to modulate expression or activity of a polypeptide or nucleic acid of the invention, a physician, veterinarian, or researcher may, for example, prescribe a relatively low dose at first, subsequently increasing the dose until an appropriate response is obtained. In addition, it is understood that the specific dose level for any particular animal subject will depend upon a variety of factors including the activity of the specific compound employed, the age, body  
25 weight, general health, gender, and diet of the subject, the time of administration, the route of administration, the rate of excretion, any drug combination, and the degree of expression or activity to be modulated.

Nucleic acid molecules (e.g., SKN-1 DNA) of the invention can be inserted into vectors and used as gene therapy vectors. Gene therapy vectors can be delivered to a subject by, for  
30 example, intravenous injection, local administration (see, e.g., U.S. Patent 5,328,470) or by stereotactic injection (see, e.g., Chen *et al.* (1994) *Proc. Natl. Acad. Sci. USA* 91:3054-3057).

The pharmaceutical preparation of the gene therapy vector can include the gene therapy vector in an acceptable diluent, or can comprise a slow release matrix in which the gene delivery vehicle is imbedded. Alternatively, where the complete gene delivery vector can be produced intact from recombinant cells, e.g., retroviral vectors, the pharmaceutical preparation can include one or  
5 more cells which produce the gene delivery system.

The pharmaceutical compositions can be included in a container, pack, or dispenser together with instructions for administration.

The invention is illustrated in part by the following examples, which are not to be taken as limiting the invention in any way.

10 Example 1: SKN-1 Links *C. elegans* Mesendodermal Specification to a Conserved Oxidative Stress Response

***C. elegans* strains and bioinformatics**

15 Strains were maintained at 20°C unless otherwise noted, using standard methods (Brenner, Genetics 77: 71-94 (1974)). Alleles used were N2 Bristol as the wild-type, and *skn-1(zu67)* and *(zu129)* (Bowerman et al. Cell 68: 1061-1075 (1992)). *C. elegans* orthologs of Nrf targets and other detoxification genes were identified by searching WORMpep or genomic databases (Sanger Centre). Predicted SKN-1 sites (Fig. 1C) 5' of their coding regions were  
20 identified with TFSEARCH (Heinemeyer et al. Nucleic Acids Res 26: 362-367 (1998)).

**Paraquat sensitivity and lifespan assays**

To assay sensitivity to paraquat, young adults were transferred from NGM agar plates into 24-well plates (6 per well) containing 0.3 µl of M9 that either did or did not contain 100mM  
25 paraquat. Worms were incubated at 20°C, and the number of dead animals was counted by the continuous absence of swimming movements and pharyngeal pumping. Lifespan assays were performed essentially as described by Hsin et al. (Nature 399: 362-366 (1999)). Animals were transferred to new plates daily and classified as dead when they did not move after repeated prodding with a pick. Animals that crawled away from the plate, exploded, or contained  
30 internally hatched worms were excluded from the analysis.

### Plasmid constructions

All PCR was performed using Pfu polymerase (Stratagene). GFP vectors pPD95.67 and pPD114.35 were obtained. A *skn-1::gfp* promoter fusion construct (SknPro::GFP; Fig. 5A) was created by ligating GFP vector pPD95.67 and an PCR-amplified 2.1 kb clone containing the promoter region and 38 amino acids from first ATG codon of the *skn-1* gene from cosmid T19E7 (sequence information about this cosmid can be accessed in WormBase under number T19E7). To generate the SKN-1::GFP translational fusion construct (Fig. 5A), the 5.7 kb EcoRI DNA fragment that rescues the maternal *skn-1* phenotype and encodes the 533 amino acid SKN-1 protein (Bowerman et al. *Cell* 68: 1061-1075 (1992)) was amplified from cosmid B0547. A ClaI site was created immediately 3' to the SKN-1 C-terminus by the Quick Change method (Stratagene), which was used for all site-directed mutagenesis. This EcoRI fragment was subcloned into pUC18 that contained the upstream 1.3 kb SphI-EcoRI fragment from SknPro::GFP (Fig. 5A). A 0.8 kb ClaI fragment that contained the GFP open reading frame (amplified from plasmid pPD114.35) was then cloned into the ClaI site to generate an in-frame exon fusion of GFP to the SKN-1 C-terminus.

The *C. elegans gcs-1* ORF (WormBase number F37B12.2) is between 45% and 54% identical to human, mouse, *Drosophila* and yeast GCS(h) (data not shown). To construct the *gcs-1::gfp* transgene, a fragment which contained 1840 bp upstream of the initiation ATG, along with sequences encoding the 17 amino terminal GCS-1 residues, was amplified by PCR from cosmid F37B12, and cloned into GFP vector pPD95.67. Promoter deletions were similarly constructed by PCR. In *gcs-1* point mutation constructs, predicted SKN-1 sites (underlined) were altered as follows: Site 1 -608 GATGACAAT to CTGCAGAAT, Site 2 -317 GATGACTTA to CTGCAGTTA, and Site 3 -121 TTTATCATC to TTTCTGCAG.

### Transgenic analyses

Transgenic strains were generated by injecting DNA into the gonad of young adult animals as described in Mello et al. (EMBO Journal 10: 3959-3970 (1991)). *gcs-1::gfp* transgene constructs (Fig. 3A) were injected at 50 ng/μl along with the *rol-6* marker (pRF4) at 100 ng/μl. Between three and six independent extrachromosomal lines were generated and analyzed for each *gcs-1::gfp* construct. To investigate GCS-1::GFP expression in the *skn-1(zu67)* background, *rol-6*-marked *gcs-1::gfp* hermaphrodites were mated with N2 males, then their

transgenic progeny were crossed with *skn-1(zu67)/DnT1* hermaphrodites, which have an *unc* phenotype. After transgenic males were successively crossed twice with *skn-1(zu67)/DnT1* hermaphrodites, *unc; rol* F3 hermaphrodite progeny were selected. From this population, *skn-1(zu67)/DnT1; gcs-1::gfp* animals were identified on the basis of their non-*unc; rol* progeny laying dead eggs. Two different *gcs-1::gfp* lines were thereby crossed into the *skn-1(zu67)* background and examined for GFP expression. DIC and fluorescence images were acquired with a Zeiss AxioSKOP2 microscope and AxioCam cooled color digital camera.

To investigate expression of *gcs-1::gfp* and mutant transgenes, worms were exposed to oxidative stress under the following conditions. For heat shock, worms cultured at 20°C were transferred onto prewarmed seeded plates and incubated at 29°C for 20 hours, then examined by fluorescence microscopy for GFP expression. *gcs-1::gfp* induction was also observed in an alternative heat treatment protocol, during which worms cultured at 20°C were transferred onto prewarmed plates and incubated at 34°C for 2 to 4 hours, then returned to 20°C and examined for GFP expression hourly during a four hour recovery period. In the experiments described in Table 2, young adults were transferred to plates that contained 1mM paraquat in the agar and maintained at 20°C for 3 days prior to analysis. In an alternative induction protocol, worms that carried *gcs-1::gfp* or the mutant transgenes shown in Figure 3A were incubated in M9 either with or without 100 mM paraquat for 30 minutes, then allowed to recover on plates for four hours. The latter procedure also resulted in induction of intestinal *gcs-1::gfp* expression by paraquat but was associated with a higher background in uninduced animals.

To create transgenic *skn-1::gfp* strains, 2.5, 10, or 50 ng/μl of transgene DNA (Fig. 5A) was injected into N2 animals along with 100 ng/μl of pRF4 to generate extrachromosomal transgenic lines. Two different extrachromosomal arrays, *Ex001* and *Ex007*, generated with 2.5 and 10 ng/μl of SKN-1::GFP, respectively, were integrated into the chromosome by UV irradiation (400J/m<sup>2</sup>) to produce the insertion strains *Is001* and *Is007*, respectively. To rescue the embryonic lethality of a *skn-1* mutation, SKN-1::GFP was injected into *skn-1(zu67)/DnT1* animals at 2.5 ng/μl with 100 ng/μl of the pRF4 marker. Rescue of maternal *skn-1* lethality was observed in some *rol*; non-*unc* progeny but not in non-*rol*; non-*unc* animals. SKN-1::GFP expression analyses shown were performed in the *Is007* strain, but essentially the same results were obtained in analyses of *Ex001*, *Ex007*, and *Is001* (data not shown). To analyze expression and localization of SKN-1::GFP in response to oxidative stress, *skn-1::gfp* transgenic worms

were treated as described above for the *gcs-1::gfp* expression studies. In addition, for exposure to sodium azide animals cultured at 20°C were placed upon a 2% agarose pad on a slide in M9 either with or without 50 mM sodium azide, then covered with a slip and examined by fluorescence microscopy. These worms were scored for presence of SKN-1::GFP in intestinal nuclei 5 minutes later. For photography, worms were immobilized either 2 mM Sodium Azide (Fig. 2, 3) or 2 mM Levamisole (Fig. 5). These treatments did not stimulate either GCS-1::GFP induction or SKN-1::GFP relocalization during the times examined (data not shown). No immobilization agent was used in the experiments shown in Tables 2 and 3. To discriminate intestinal autofluorescence from SKN-1::GFP epifluorescence, a triple band emission filter set (Chroma 61000) was used in conjunction with a narrow band excitation filter (484/14 nm). This combination allowed autofluorescence to be detected as yellow/orange fluorescence deriving from a combined green and red signal, while GFP remained green. Worms that carried *skn-1::gfp*, *gcs-1::gfp* and *gcs-1::gfp* mutant transgenes were incubated with 50 µg/ml DiI (Molecular Probes) in M9 for 3 hours at 20°C, then transferred to fresh plates for 1 hour to destain, and examined under the fluorescence microscope. The ASI chemosensory neurons were identified by according to their intensity of DiI labeling and location relative to other DiI-labeled cells.

#### DNA binding assays

Full-length SKN-1 and the SKN domain were expressed by in vitro translation (Promega) as described previously (Carroll et al., Genes Dev. 11: 2227-2238 (1997)). Oligonucleotide probes were end-labeled using Klenow and  $\alpha$ -<sup>32</sup>P-labelled dATP and CTP, then purified using QIAquick Kit (Qiagen). EMSAs were performed essentially as described in Blackwell et al. (Science 266: 621-628 (1994)), with labeled probes present at  $2.5 \times 10^{-9}$  M.

#### Constitutive and inducible Phase II detoxification gene activation by SKN-1

Vertebrate Nrf proteins induce expression of Phase II detoxification enzyme genes by binding to the characteristic antioxidant response element (ARE) in their promoters (Fig. 1C) (Hayes et al. Cancer Lett 174: 103-113 (2001)). A search was performed for SKN-1 binding sites within the predicted promoters of *C. elegans* orthologs of these oxidative stress resistance genes. The SKN-1 binding site preference and the ARE are distinct but not mutually exclusive (Fig. 1C). A predicted SKN-1 site should appear randomly every 2048 bp, but between two and

four SKN-1 sites are present within 1 kb upstream of multiple *C. elegans* genes that encode predicted Phase II detoxification enzymes, including  $\gamma$ -glutamine cysteine synthase heavy chain (GCS(h)), glutathione synthetase, and four glutathione S-transferase (GST) isoforms (Table 1). In vertebrates each of these genes is activated by Nrf proteins (Id.). SKN-1 sites or variants that differ at only one AT-rich region position are similarly present 5' of the Nrf target *NADH* quinone oxidoreductase, the catalase *ctl-1*, and superoxide dismutases (*sod-1*, -2 and -3) (see Table 1, below).

Table 1. Predicted SKN-1 binding sites upstream of *C. elegans* oxidative stress resistance genes

Enzymes	Gene or ORF	Location <sup>a</sup>	Direction	Sequence
$\gamma$ -glutamyl-cysteine synthetase heavy chain (GCS(h))	<i>gcs-1</i>	-121	→	TTTATCAT
		-316	←	ATGACTTA
Glutathione synthetase	M176.2	-607	←	ATGACAAT
		-137	→	TTTGTCAT
		-169	←	ATGACAAA
		-243	→	TTTATCAT
		-378	←	ATGATTTT
		-469	→	GTTATCAT
NADH quinone oxidoreductase	F39B2.3	-518	←	ATGACAAT
Glutathione S-transferase	R03D7.6	-149	←	ATGACAAT
		-282	←	ATGATTTT
		-302	←	ATGACATT
		-947	←	ATGATTTT
	F35E8.8	-94	←	ATGACAAT
		-240	←	ATGATAAT
Superoxide dismutase	F11G11.2	-133	←	ATGACAAA
		-391	→	CTTATCAT
	K08F4.7	-83	←	ATGACATT
		-157	→	TTTGTCAT
	<i>sod-1</i>	-64	→	ATAATCAT
	<i>sod-2</i>	-191	→	TGTATCAT
		-363	←	ATGACAAT
		-959	→	AGAATCAT
		-980	→	AGAATCAT
Catalase	<i>sod-3</i>	-287	→	TAAATCAT
	<i>ctl-1</i>	-380	→	ATGATCAT
		-978	→	GTCATCAT
		-997	→	CTTATCAT



The SKN-1 binding consensus is shown in Figure 1C. <sup>a</sup>The A within the translation initiation codon is designated as base 1.

5 Presence of SKN-1 site clusters upstream of multiple *C. elegans* Phase II detoxification genes is consistent with SKN-1 functioning analogously to Nrf proteins. To confirm this, whether SKN-1 is required to express the Phase II gene *gcs-1* (Table 1) was investigated. *gcs-1* is the *C. elegans* ortholog of *GCS(h)*, a representative and well-characterized Nrf protein target gene that in yeast is regulated by Yap1p and Pap1p (Id., and Toone et al., Curr Opin Genet Dev 9: 55-61 (1999)). The GCS(h) enzyme is important for oxidative stress resistance because it is rate-limiting for glutathione synthesis.

10 *gcs-1* expression in *C. elegans* was investigated using a transgene that included the predicted *gcs-1* promoter, along with the 17 amino terminal GCS-1 amino acids fused to green fluorescent protein (GFP). This promoter segment contained three consensus SKN-1 binding sites, and corresponded to the intervening sequence between *gcs-1* and the nearest upstream gene (data not shown). With this strategy, it was possible to analyze *gcs-1* expression independently of GCS-1 protein stability. In a wild-type background, during larval and adult stages GCS-1::GFP was readily detectable in the pharynx, and in nearby cells that appeared to be neurons (Figs. 2A and B). By soaking *gcs-1::gfp* lines in DiI, a dye that fills amphid sensory neurons (Herman et al., Nature 348:169-171 (1990)), it was determined that two GCS-1::GFP-expressing cells located adjacent to the posterior pharynx correspond to the ASI chemosensory neurons (Fig. 2C and 2D), which prevent constitutive entry into the dauer diapause state (Ren et al. 1996; Schackwitz et al. 1996). GCS-1::GFP expression was also apparent anteriorly and posteriorly in the intestine (Figs. 2A and 2B).

25 In vertebrates, oxidative stress induces Phase II gene expression through an Nrf2-dependent pathway in the intestine and liver (Itoh et al. 1997; Hayes and McMahon 2001). Similarly, stimuli that cause oxidative stress dramatically increased GCS-1::GFP expression in the *C. elegans* intestine (Figs. 2E and 2F, and see Table 2, below). This response was triggered by both heat and the herbicide paraquat (methyl viologen), which generates intracellular superoxide anions. To investigate the involvement of *skn-1* in *gcs-1* expression, we introduced the *gcs-1::gfp* transgene into the *skn-1(zu67)* background, the *skn-1* allele that is associated with the most severe embryonic phenotype (Bowerman et al. Cell 68: 1061-1075 (1992)). Under both normal and oxidative stress conditions, in *skn-1(zu67)* homozygotes GCS-1::GFP was apparent

at wild-type levels in the pharynx, but was otherwise undetectable (Figs. 2G-2L), indicating that *skn-1* is essential for both constitutive and inducible *gcs-1::gfp* expression outside of the pharynx.

Promoter mutagenesis identified discrete elements that are required for these *skn-1*-dependent and -independent *gcs-1* expression patterns. Pharyngeal GCS-1::GFP expression was abolished by removal of the distal *gcs-1* promoter region (*gcsΔ2::gfp*; Figs. 3A and 3B), which lacks SKN-1 binding sites but contains consensus sites for the pharyngeal transcription factors PEB-1 and PHA-4 (Thatcher et al. *Dev Biol* 229: 480-493 (2001); Gaudet et al., *Science* 295: 821-825 (2002)) (data not shown). The remaining proximal 682 bp of the *gcs-1* promoter included the three predicted SKN-1 binding sites, and was sufficient for appropriate GCS-1::GFP expression in the intestine and ASI neurons (*gcsΔ2::gfp*, Figs. 3A, 3B, Table 2). Constitutive and stress-induced GCS-1::GFP expression within the intestine and ASI neurons did not require SKN-1 binding sites 1 or 2 individually, but was abolished by alteration of site 3 (*gcs2;mut3::gfp*; Figs. 3A and 3B).

**Table 2.** Induction of GCS-1::GFP expression in the intestine by oxidative stress

Inducer	<i>gcs-1::gfp</i>				<i>gcsΔ2::gfp</i>			
	low	medium	high	N	low	medium	high	N
Control	90.8%	7.9%	1.3%	76	88.2%	10.3%	1.5%	68
Heat Shock	10.5%	72.4%	17.1%	76	0.0%	14.0%	86.0%	86
Paraquat	14.5%	67.1%	18.4%	76	21.3%	65.6%	13.1%	61

A representative set of experiments involving a mixed population of L2-young adult worms is shown, from which percentages of animals in each expression category are listed. Induction of GCS-1::GFP expression was comparable among the different developmental stages analyzed. "Low" refers to animals similar to that in Fig. 2A, in which intestinal GCS-1::GFP was apparent at modest levels anteriorly, or anteriorly and posteriorly. "High" indicates that a GCS-1::GFP was present at high levels anteriorly and detectable throughout most of the intestine, as in Fig. 2F. "Medium" refers to animals in which GCS-1::GFP was present at high levels anteriorly as in Fig. 2F and possibly posteriorly, but was not detected in between. N indicates numbers of animals analyzed from each transgenic strain.

Remarkably, SKN-1 binding site 3 is located within a 42 bp *gcs-1* promoter element that is similar to a composite motif through which SKN-1 activates *med-1* and *med-2* in the embryo (Figs. 1A and 3C) (Maduro et al. Mol. Cell 7: 475-485 (2001)). The conservation between these *gcs-1* and *med* promoter elements is particularly striking because they are located at identical distances from their respective translation starts, but contain different numbers of SKN-1 sites (Figs. 3C). In an electrophoretic mobility shift assay (EMSA), full-length SKN-1 and the 85 amino acid SKN-1 DNA binding domain (SKN Domain) (Blackwell et al. Science 266: 621-628 (1994)) each bound sequence-specifically to SKN-1 binding site 3 in the context of this *gcs-1* promoter element (Fig. 4). These SKN-1 proteins bound with high affinity to an oligonucleotide that corresponds to this composite element (Wild type; Figs. 4A and 4B, lanes 2-5), but not to an analogous probe in which SKN-1 site 3 had been altered as in the inactive *gcsΔ2; mut3::gfp* transgene (Fig. 3A; Mutant, Figs. 4A and 4B, lanes 7-10). Binding of these SKN-1 proteins to the Wild type probe was also competed much more effectively by unlabeled Wild type than Mutant DNA (Figs. 4A and B, lanes 11-20). Further supporting the importance of this *gcs-1* promoter element, a 163 bp fragment that includes it provides significant GCS-1::GFP expression in the intestine, but 5' truncation within this sequence inactivates the promoter (*gcsΔ4::gfp* and *gcsΔ5::gfp*, Fig. 3A). It is evident that binding of SKN-1 to site 3 is required for *gcs-1* expression in the intestine and ASI neurons.

#### SKN-1 expression and accumulation in intestinal nuclei in response to oxidative stress

To determine whether SKN-1 is present in tissues where it is required for *gcs-1::gfp* expression, expression of a transgene in which GFP is fused to the C-terminus of full-length SKN-1 (SKN-1::GFP; Fig. 5A) was analyzed. Although maternal *skn-1::gfp* expression was not readily detectable because of germline transgene silencing (Kelly et al., Genetics 146:227-238 (1997)), at a low frequency this transgene rescued the embryonic defect in *skn-1(zu67)* homozygotes (data not shown), indicating that this SKN-1::GFP fusion protein is functional.

In the embryo, antibody staining previously revealed presence of maternal SKN-1 in nuclei through the eight-cell stage, then detected zygotically expressed SKN-1 in only about 15% of late-stage embryos that had ceased dividing (Bowerman et al. Cell 74: 443-452 (1993)).

Nuclear SKN-1::GFP was uniformly detected in intestinal precursors beginning at the 50-100 cell stage (Fig. 5B), then in both the intestine and hypodermis (Fig. 5C), indicating that SKN-1 is expressed zygotically earlier than it is detectable by antibody staining. In late-stage embryos

SKN-1::GFP was also present in intestinal nuclei but not in the hypodermis (Fig. 5D), suggesting that hypodermal *skn-1* expression may be maintained by a region located outside of this transgene.

In contrast to the embryo, in larvae and young adults SKN-1::GFP was usually present at very low levels in intestinal nuclei (Fig. 5F, and see Table 3, below). SKN-1::GFP was readily detectable in the ASI neurons, where *gcs-1::gfp* was constitutively expressed (Figs. 5E and 5F), but not in other cells in the head where GCS-1::GFP expression appeared to be *skn-1*-dependent (Figs. 2B and 2H). The latter *skn-1* dependence might be indirect, or derived from low level SKN-1 expression or distant *skn-1* regulatory regions. The finding that SKN-1::GFP is present at only modest levels in intestinal nuclei raises the question of how oxidative stress induces *skn-1*-dependent intestinal *gcs-1* expression (Figs. 2, 3, and Table 2, above). In cultured mammalian cells, Nrf2 is stabilized and relocalized from the cytoplasm to the nucleus in response to oxidative stress (Itoh et al. *Genes Dev* 13: 76-86 (1999); Sekhar et al. *Oncogene* 21: 6829-6834 (2002); Nguyen et al. *J Biol Chem* 278: 4536-4541 (2003); and Stewart et al. *J Biol Chem* 278: 2396-2402 (2003)). A promoter fusion transgene in which only the SKN-1 amino terminus was linked to GFP (SknPro::GFP, Fig. 5A) was constitutively expressed at high levels in all intestinal cells (data not shown), suggesting that SKN-1 expression or localization might also be regulated post-transcriptionally by oxidative stress.

After exposure to either paraquat or heat, neither the location nor intensity of SKN-1::GFP was detectably altered in the ASI neurons, but in a high percentage of animals elevated levels of SKN-1::GFP appeared in intestinal cell nuclei, particularly anteriorly and posteriorly where GCS-1::GFP is most robustly expressed (Figs. 5F and 5G, Table 3). SKN-1::GFP accumulated in intestinal nuclei within 5 min. after treatment with 50 mM sodium azide (Table 3), which induces oxidative stress by blocking mitochondrial electron transport. The rapidity of this last response indicates that in the intestine SKN-1 is constitutively present, but may be diffuse within the cytoplasm and masked by autofluorescence. This accumulation of SKN-1::GFP in intestinal nuclei in response to oxidative stress remarkably parallels the *skn-1*-dependent induction of GCS-1::GFP under similar conditions, supporting the model that SKN-1 activates intestinal *gcs-1* expression directly.

**Table 3.** Accumulation of SKN-1::GFP in intestinal nuclei in response to oxidative stress

Inducer	low	medium	high	N
Control	78.9%	14.5%	6.6%	76
Heat	5.6%	11.9%	82.5%	143
Paraquat	53.1%	43.8%	3.1%	64
M9, 5 min.	74.7%	17.6%	7.7%	91
50 mM Sodium Azide, 5 min.	0.8%	44.2%	55.0%	120

Mixed-stage L2-young adult transgenic worms were exposed to the indicated conditions. A representative set of experiments is shown, from which percentages of animals in each category are listed. SKN-1 localization patterns did not differ significantly among the different developmental stages examined. M9 refers to the control incubation for the sodium azide experiment. In some animals treated with sodium azide, high levels of nuclear SKN-1::GFP appeared in less than 1 minute (data not shown). "Low" refers to animals in which SKN-1::GFP was barely detectable in all intestinal nuclei, as shown in Fig. 4F. "High" indicates that a very strong SKN-1::GFP signal was present in all intestinal nuclei, as in Fig. 4G. "Medium" refers to animals in which nuclear SKN-1::GFP was present at high levels anteriorly or anteriorly and posteriorly, but was barely detectable midway through the intestine. N indicates numbers of animals analyzed in each category.

### **skn-1 required for oxidative stress resistance and normal longevity**

Whether *skn-1* mutants are abnormally sensitive to oxidative stress was investigated. *skn-1(zu67)* homozygotes produce normal numbers of offspring with normal timing, and as young adults are not obviously distinguishable in morphology from wild-type (data not shown). Two different *skn-1* mutant alleles were associated with markedly decreased survival in the presence of paraquat however, indicating that *skn-1* mutants are sensitive to oxidative stress (Fig. 6A).

Further, whether *skn-1* homozygotes live as long as wild type was investigated. Both the mean and maximum lifespans of *skn-1(zu67)* and *skn-1(zu129)* homozygotes were reduced by 25-30% (Fig. 6B), indicating that SKN-1 is required for normal longevity.

### **A conserved postembryonic function for SKN-1 in oxidative stress resistance**

The *C. elegans* developmental specification protein SKN-1 also mediates a conserved response to oxidative stress. SKN-1 functions similarly to bZIP proteins that regulate Phase II

detoxification genes in vertebrates (Nrf1, 2) and yeast (Yap1p, Pap1p). SKN-1 activates a conserved Phase II gene in the intestine and ASI neurons (Figs. 2, 3, 5), SKN-1 binding sites flank *C. elegans* orthologs of additional Nrf target genes (Table 1), and *skn-1* mutants are sensitive to oxidative stress (Fig. 6A). The accumulation of SKN-1 in intestinal nuclei in response to oxidative stress (Fig. 5G, and Table 3) may parallel nuclear accumulation of Nrf proteins, Yap1p, and Pap1p, under these conditions (Itoh et al. *Genes Dev* 13: 76-86 (1999); Toone et al. *Oncogene* 20: 2336-2346 (2001); and Dclanay et al., *Cell* 111: 471-81 (2002)). The intestinal abnormalities in *skn-1(zu67)/nDf41* larvae (Bowerman et al. *Cell* 68: 1061-1075 (1992)) could involve oxidative stress, because 10-20% of *gcs-1(RNAi)* animals also die as larvae with abnormal intestines (data not shown).

These parallels between SKN-1 and Nrf proteins are surprising because the mechanism through which SKN-1 binds DNA is both unique and highly divergent (Blackwell et al. *Science* 266: 621-628 (1994)). SKN-1 and Nrf proteins are most similar within the 14 amino acid DIDLID transactivation element (Fig. 1B) (Walker et al. *J. Biol. Chem.* 275: 22166-22171 (2000)).

While *gcs-1* expression in the intestine is induced by SKN-1 in response to stress, presence of nuclear SKN-1 allows *gcs-1* to be expressed constitutively in the ASI neurons, and *gcs-1* expression is *skn-1*-independent in the pharynx (Fig. 2). In metazoans, Phase II genes thus can be activated through distinct pathways that may be important for functions of different tissues. For example, the finding that *skn-1* functions constitutively in the ASI neurons, which inhibit dauer entry, suggests that although *skn-1(zu67)* homozygotes can enter the dauer stage (data not shown), *skn-1* or oxidative stress might influence regulation of this process.

The lifespan reduction that observed in *skn-1* mutants (25-30%, Fig. 6B) is comparable to that reported in *daf-16* mutants (20%) (Kenyon et al. *Nature* 366: 461-464 (1993); and Lee et al. *Curr Biol* 11: 1950-1957 (2001)). In *C. elegans*, aging involves pleiotropic changes that vary among individuals, and mutations that influence lifespan may affect aging of some tissues more than others (Garigan et al. *Genetics* 161: 1101-1112 (2002); and Herndon et al. *Nature* 419: 808-814 (2002). Just before death the anterior intestine and posterior pharynx degenerated more frequently in *skn-1* animals than wild type (data not shown), a finding that may reflect aging but does not exclude the possibility of an additional defect. At one week after hatching, small cavities and apparent yolk droplets appeared in the heads of many *skn-1* but not wild type

animals (data not shown). These changes in the head region are typical of aging *C. elegans* (Garigan et al. *Genetics* **161**: 1101-1112 (2002); Herndon et al. *Nature* **419**: 808-814), suggesting that *skn-1* mutants age prematurely. Some mechanisms that regulate *C. elegans* lifespan have been shown to influence lifespan in higher metazoans (Clancy et al. *Science* **292**: 104-106 (2001); Finch et al. *Annu Rev Genomics Nature* **421**: 182-187 (2003)). The observation that normal *C. elegans* longevity requires *skn-1* is consistent with other associations between oxidative stress resistance and lifespan, and suggests that the conserved oxidative stress resistance pathway regulated by SKN-1 might influence longevity in other species.

#### OTHER EMBODIMENTS

It is to be understood that while the invention has been described in conjunction with the detailed description thereof, the foregoing description is intended to illustrate and not limit the scope of the invention, which is defined by the scope of the appended claims. Other aspects, advantages, and modifications are within the scope of the following claims.

**WHAT IS CLAIMED IS:**

1. A method for determining whether a test compound is a candidate SKN-1-mediated oxidative stress response-activating compound, comprising:

(a) providing a nematode capable of expressing a SKN-1 polypeptide and containing at least one transgene comprising:

(i) an oxidative stress resistance gene promoter operably linked to

(ii) a reporter gene; and

(b) contacting the nematode with the test compound; and

(c) determining whether expression of the transgene is increased, wherein an increase in expression of the transgene indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-activating compound.

2. A method for determining whether a test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound, comprising:

(a) providing a nematode capable of expressing a SKN-1 polypeptide and containing at least one transgene comprising:

(i) an oxidative stress resistance gene promoter operably linked to

(ii) a reporter gene;

(b) contacting the nematode with the test compound; and

(c) before, during, or after step (b), subjecting the nematode to conditions that activate the SKN-1-mediated oxidative stress response in the absence of the test compound; and

(d) determining whether expression of the transgene is decreased or unchanged, wherein decreased or unchanged expression of the transgene indicates that the test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting compound.

3. A method for determining whether a test compound is a candidate SKN-1-mediated oxidative stress response-activating compound, comprising:

(a) providing a nematode containing a transgene encoding a SKN-1 fusion protein, wherein the transgene comprises:

(i) a SKN-1 DNA operably linked to

(ii) a reporter gene;



- 32 (b) contacting the nematode with the test compound; and  
33 (c) determining whether the SKN-1 fusion protein accumulates in nuclei in the  
34 nematode, wherein increased accumulation indicates that the test compound is a candidate  
35 SKN-1-mediated oxidative stress response-activating compound.  
36
- 37 4. A method for determining whether a test compound is a candidate SKN-1-mediated  
38 oxidative stress response-inhibiting compound, comprising:  
39 (a) providing a nematode containing a transgene encoding a SKN-1 fusion protein,  
40 wherein the transgene comprises:  
41 (i) a SKN-1 DNA operably linked to  
42 (ii) a reporter gene;  
43 (b) contacting the nematode with the test compound;  
44 (c) before or during step (b), subjecting the nematode to conditions that activate the  
45 SKN-1-mediated oxidative stress response in the absence of the test compound; and  
46 (d) determining whether the SKN-1 fusion protein accumulates in nuclei in the  
47 nematode, wherein decreased or unchanged accumulation of the transgene indicates that the  
48 test compound is a candidate SKN-1-mediated oxidative stress response-inhibiting  
49 compound.  
50
- 51 5. The method of claim 1 or 3, further comprising the step of:  
52 (d) contacting a second nematode, or a cultured nematode cell, or cultured  
53 mammalian cell with the candidate compound to determine whether the candidate compound  
54 increases oxidative stress resistance, relative to the oxidative stress resistance of the second  
55 nematode, the cultured nematode cell, or the cultured mammalian cell not contacted with the  
56 candidate compound, wherein a candidate compound that increases oxidative stress  
57 resistance is an oxidative stress response-activating agent.  
58
- 59 6. The method of claim 2 or 4, further comprising the step of:  
60 (e) contacting a second nematode, or a cultured nematode cell, or cultured  
61 mammalian cell with the candidate compound to determine whether the candidate compound  
62 decreases oxidative stress resistance, relative to the oxidative stress resistance of the second

63 nematode, the cultured nematode cell, or the cultured mammalian cell not contacted with the  
64 candidate compound, wherein a candidate compound that decreases oxidative stress  
65 resistance is an oxidative stress response-inhibiting agent.  
66

67 7. The method of claim 1 or 2, wherein the promoter is a promoter of a gene encoding a  
68 protein selected from the group consisting of:  $\gamma$ -glutamine cysteine synthase heavy chain,  
69 glutathione synthetase, NADH quinone oxidoreductase, superoxide dismutase, catalase, and  
70 glutathione S-transferase.  
71

72 8. The method of claim 1, 2, 3, or 4, wherein the reporter gene is a gene encoding a protein  
73 selected from the group consisting of: green fluorescent protein, chloramphenicol acetyl  
74 transferase,  $\beta$  glucuronidase, and luciferase.  
75

76 9. The method of claim 1 or 2, wherein the nematode in step (a) is *Caenorhabditis elegans*.  
77

78 10. The method of claim 3 or 4, wherein the nematode in step (a) is *Caenorhabditis elegans*.  
79

80 11. A compound capable of activating a SKN-1-mediated oxidative stress response isolated  
81 by the method of claim 1 or 3.  
82

83 12. A compound capable of inhibiting a SKN-1-mediated oxidative stress response isolated  
84 by the method of claim 2 or 4.  
85

86 13. An oxidative stress response-activating agent isolated by the method of claim 5.  
87

88 14. An oxidative stress response-inhibiting agent isolated by the method of claim 6.  
89

90 15. The method of claim 1, further comprising the steps of:  
91

92 (c) providing a nematode not capable of expressing a SKN-1 polypeptide and  
93 containing at least one transgene comprising:

(i) an oxidative stress resistance gene promoter operably linked to

94 (ii) a reporter gene; and  
95 (d) contacting the nematode with the test compound, wherein no increase in  
96 expression of the transgene following step (d) indicates that the candidate compound is an  
97 oxidative stress response-activating agent.  
98

99 16. A method for determining whether a test compound is a candidate compound capable of  
100 inhibiting a SKN-1-mediated oxidative stress response, comprising:

101 (a) contacting a SKN-1 polypeptide or SKN-1 DNA with a test compound ; and  
102 (b) detecting interaction of the test compound with the SKN-1 polypeptide SKN-1  
103 DNA, wherein an interaction indicates that the test compound is a candidate compound  
104 capable of inhibiting a SKN-1-mediated oxidative stress response.  
105  
106

107 17. A method for determining whether a test compound is a candidate compound capable of  
108 inhibiting a SKN-1-mediated oxidative stress response, comprising:

109 (a) providing a SKN-1 polypeptide or fragment thereof and an oxidative stress  
110 resistance gene encoding  $\gamma$ -glutamine cysteine synthase heavy chain, glutathione synthetase,  
111 NADH quinone oxidoreductase, superoxide dismutase, catalase, or glutathione S-transferase,  
112 or SKN-1 polypeptide-binding fragments thereof;

113 (b) contacting the SKN-1 polypeptide or fragment thereof and the oxidative stress  
114 resistance gene or SKN-1 polypeptide-binding fragment thereof with a test compound; and

115 (c) determining whether the SKN-1 polypeptide or fragment thereof and the oxidative  
116 stress resistance gene or SKN-1 polypeptide-binding fragment thereof interact in the  
117 presence of the test compound, wherein a decrease in interaction indicates that the test  
118 compound is a candidate compound capable of inhibiting a SKN-1-mediated oxidative stress  
119 response.  
120

**ABSTRACT**

The present invention relates, in part, to the *C. elegans* SKN-1 gene and protein (a transcription factor), and target genes thereof. The invention includes various therapeutic methods and screening methods for identifying antioxidants.

5

20669657.doc

Matter No.: 10861-032P01  
 Applicant(s):  
 SKN-1 GENE AND PROTEIN

Page 1 of 31

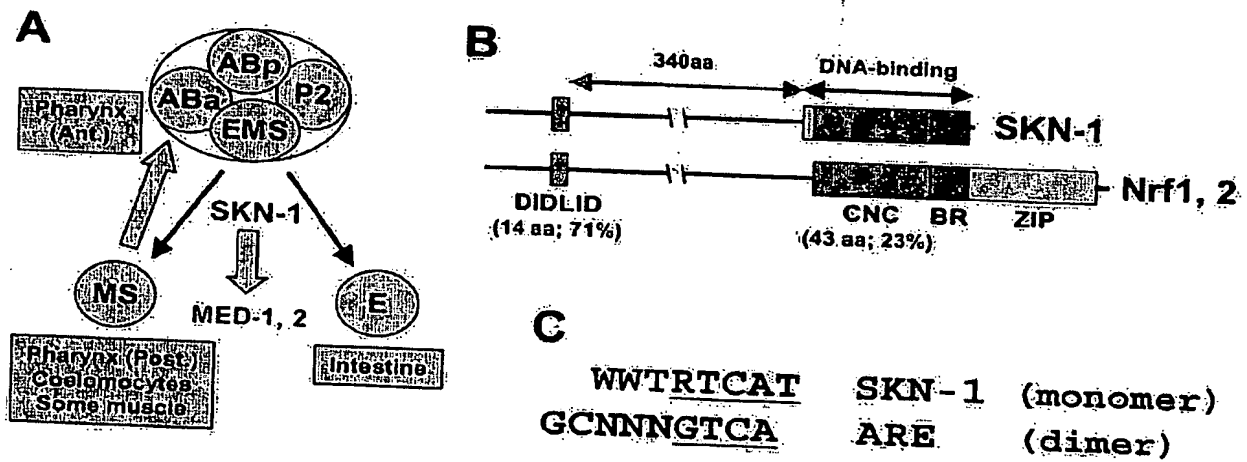
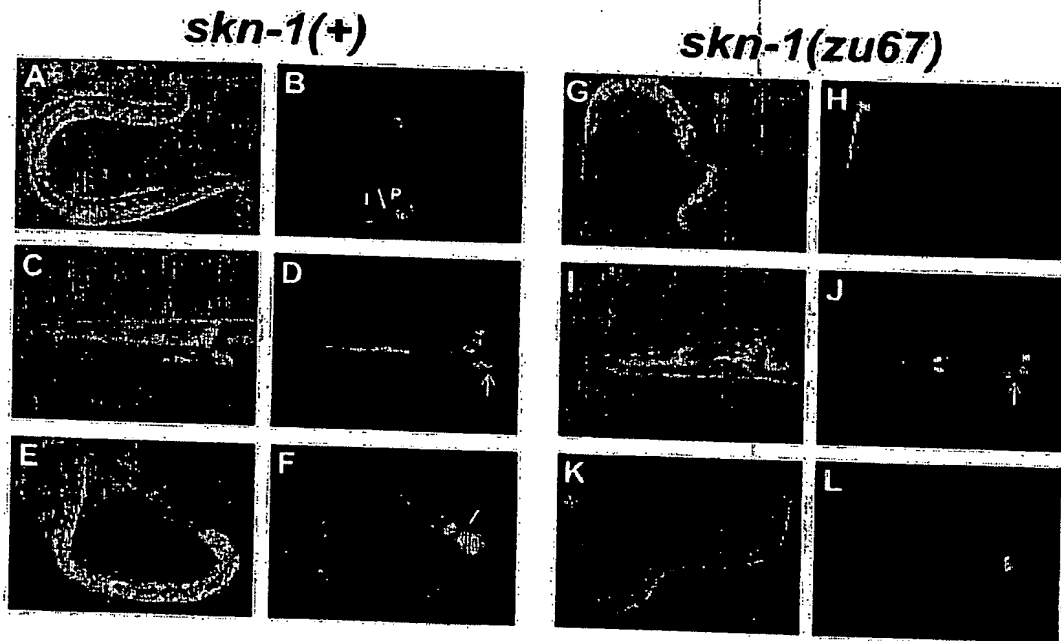


Fig. 1A-1C

Matter No.: 10861-032P01  
Applicant(s):  
SKN-1 GENE AND PROTEIN

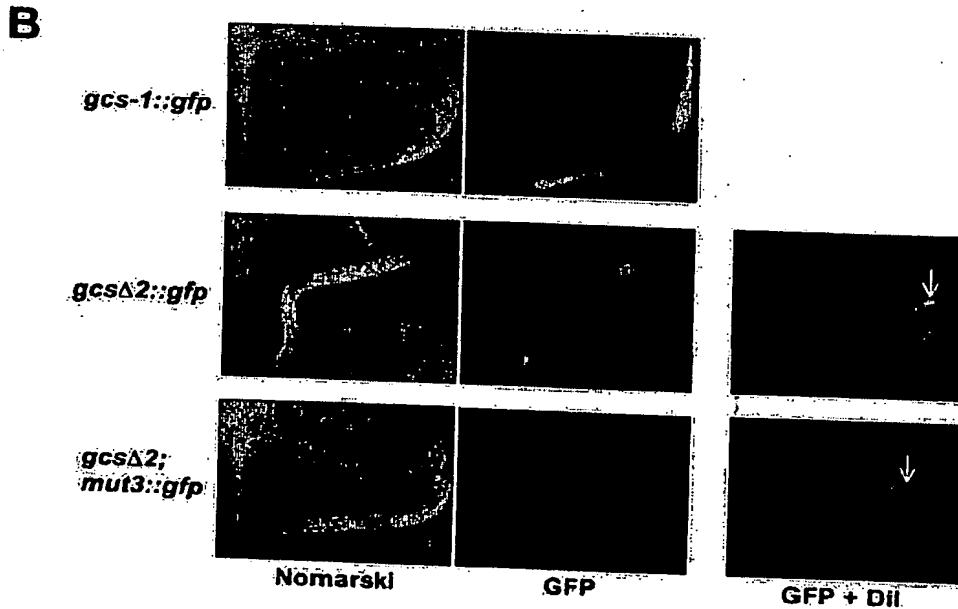
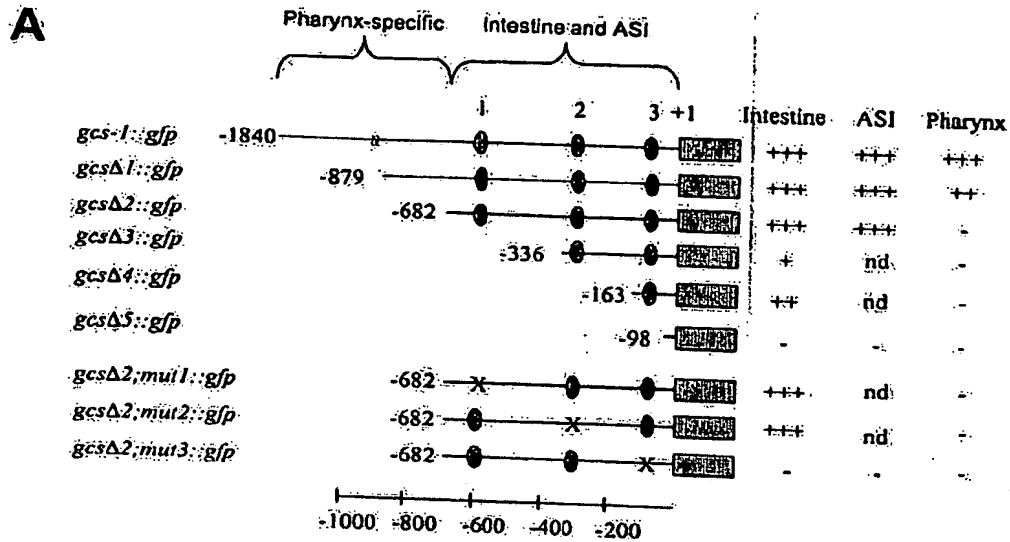
Page 2 of 31



**Fig. 2A-2L**

Matter No.: 10861-032P01  
 Applicant(s):  
 SKN-1 GENE AND PROTEIN

Page 3 of 31



**C**

<i>gcs-1</i>	-124	CA-CTTTATCATCATGA-GATTTAATCTTTCCTTTTGAATTTTCT-83
<i>med-1</i>	-127	CACCTCTGTCATCATGAATTTTTCGAG-CATTATCATCATTTCT-83
<i>med-2</i>	-127	CACCTCTGTCATCATGAATTTTTCGAG-CATTATCATCATTTCT-83

Fig. 3A-3C

Matter No.: 10861-032P01  
Applicant(s):  
SKN-1 GENE AND PROTEIN

Page 4 of 31

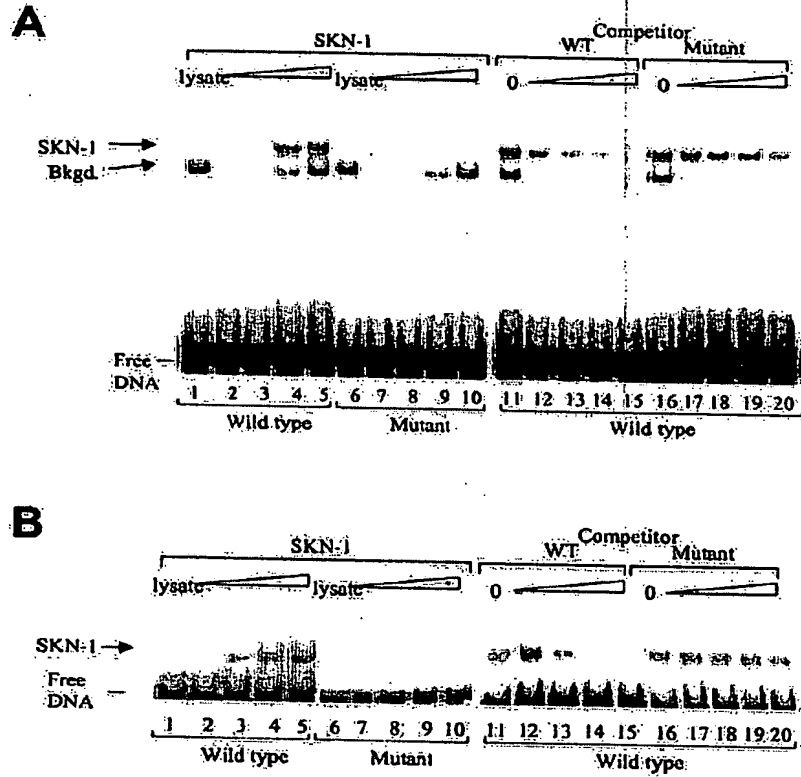


Fig. 4A-4B



Matter No.: 10861-032P01  
Applicant(s):  
SKN-1 GENE AND PROTEIN

Page 5 of 31

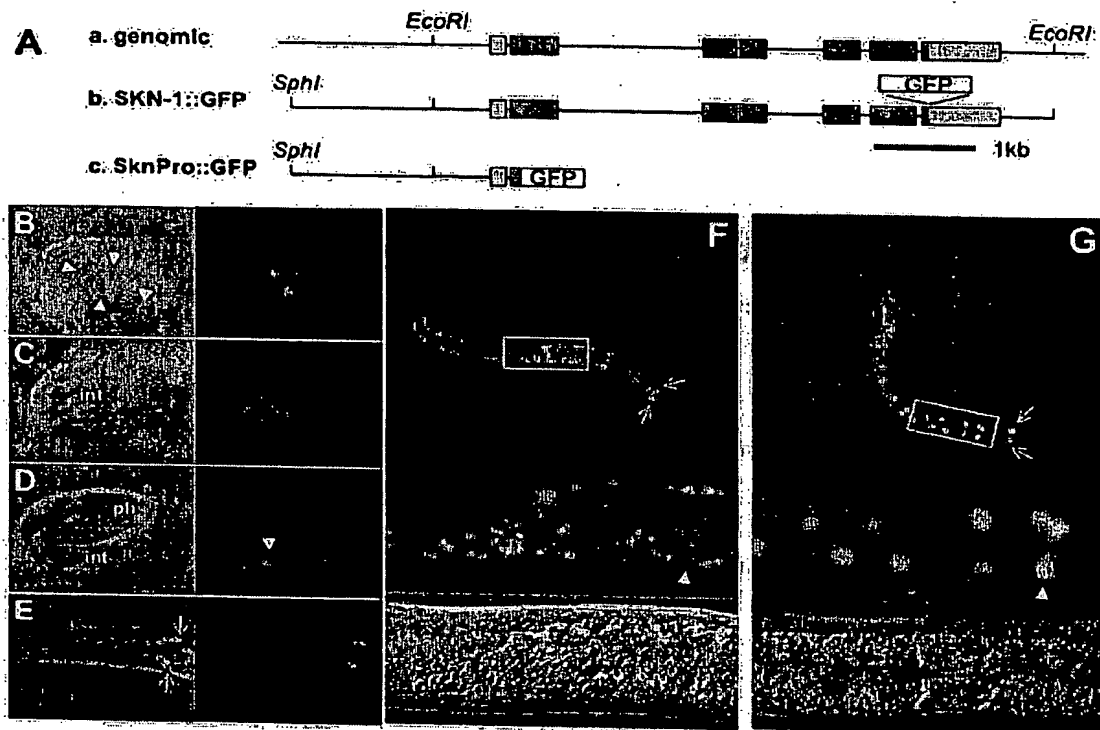


Fig. 5A-5G

Matter No.: 10861-032P01  
Applicant(s):  
SKN-1 GENE AND PROTEIN

Page 6 of 31

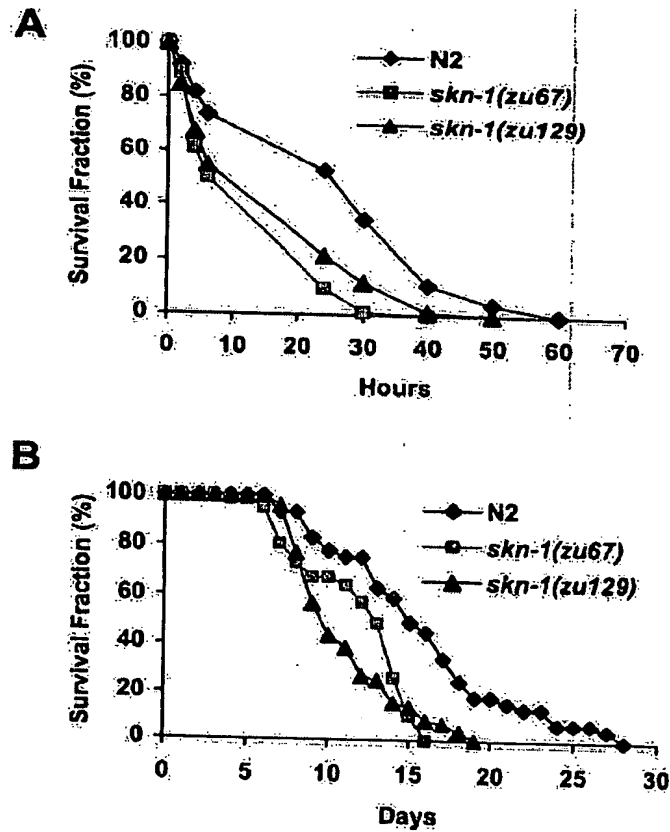
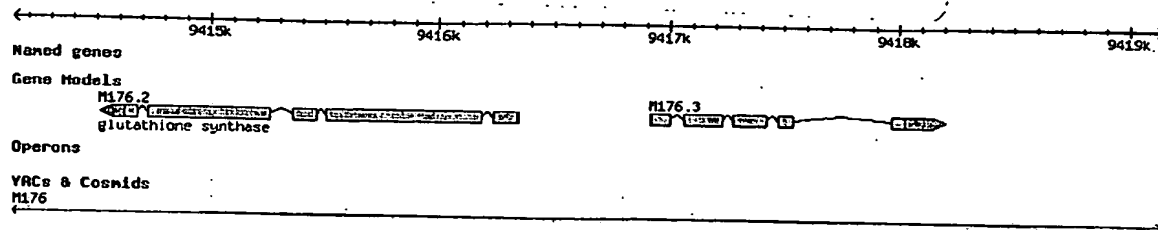


Fig. 6A-6B



The M176.2 gene is located on chromosome II. Regulatory sequences can be found e.g., in the region between 9416340 and 9415915. An exemplary sequence of this region is as follows:

```

GACAATTATCGATTAATAAAAGTTTAAACAGACACGAGAAATTAAATATAAAAAATTGAATTGTTTATTT
GTTGTTTTGTGTGTAGAAAAATAATTTTGATAGAAACAAAAATTAGCGTAAAATAAATAGCTAGCGCAA
TACTCGTGCACGAGATGTGCGCCAGCAGCTCCTTGACGCAAAACGTGACGTTTAGCACCAAATGATTTT
                                     -378
TGCTCTTTGAGTTCTTTGTTTTCGGGAGCAAATTCATGCCAATCCCTTTCTTTTTTCAAATTTTCCTG
TTAAATTCATGTAATAACTATTATTCATGTCAATTACAACAAATAAGCATCCAAGATTTTATCATAAACT
                                     -243
CGTTCAAACCTCCTTTTACCACTCGAAAAGCAATATCTCCGACTTCCTTCAAAGAGAAATGATGACAAAA
                                     -169
CATAGAAACCTCACGTTATACGTTTTGTCATCACGATTCAGTGCTCACTTTTCTCATTTTCATTCTCGCT
                                     -137
TAATTTTCATTTTTGTCACTCTCGCGTCATGTTTTGCATTTTTCGAAAGCATTATTTTAAAACTGAAAAAA
TAATTCGTAATTTTTCAAGAAATGGCT

```

FIG. 7

Matter No.: 10861-032P01  
 Applicant(s):  
 SKN-1 GENE AND PROTEIN

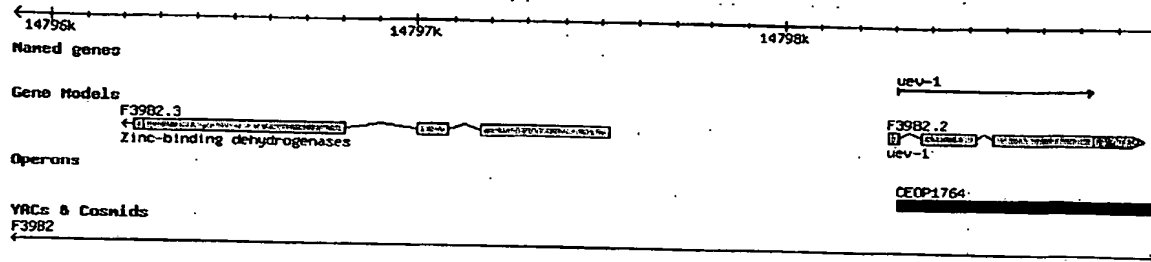
Page 8 of 31

Exemplary M176.1 sequences:

Amino Acid Sequence							
MAQKDDRILL	LNAPRLPLED	DKLNELTADL	HDWAHANGLV	MRLSTDKLSS	EVCQTTPPLT	LPSPFPKNVF	EEAVHIQNL
ASLYHFIAYE	FDFLIDIHKN	VVKTDDEFTRN	MVEILKKVKA	QGLKQPVTLA	IQRSDYCHK	DQYSAEYGLK	QIEINNIASS
MGHALRLTE	WHIRVLKALN	ISDDVIQRAI	PENKPIPMIA	EALFKAWSHF	SNPAAVVLVV	VENVNQNOID	QRHVEYELEK
LGVPMTCIIR	RNLTQCYEQL	SLNDRSDLMI	DGRQVAIVYF	RAGYSPDHYP	STKEWEARER	MELSTAIKTP	WIGLQVANTK
KTOQVLSEDE	VLERFIGKPR	EARDIRASFA	GMWALENTDE	VTMKVVAGAQ	KHPEAFVLKP	QTEGGAALHT	GDEMVOMLRE
LPEEERGAFI	LMEKLKPMII	ENYLVLAKKP	ITFAKAVSEL	GVYGYAFGRK	DAPELKTAGH	LLRTKPESTA	MGGVAAGHAV
VDTPFLYEFI							

Spliced mRNA							
aaagaATGGCT	CAAAAAGATG	ACCGGATTTT	GCTGTTGAAT	GCTCCAAGGC	TCCCGCTCGA	AGATGATAAG	CTCAACGAGC
TCACCGCTGA	TCTTCACGAT	TGGGCTCATG	CTAATGGGCT	TGTCATGCGT	CTATCAACCG	ACAAGTTGAG	CAGCGAAGTT
TGTCAAACCTA	CTCCATTAAC	ACTTCTTCCA	TCTCCATTCC	CGAAAAATGT	TTTTGAAGAA	GCAGTTCATA	TTCAGAACCT
TTTCGCAAGT	CTTTATCACT	TCATAGCTTA	TGAATTTGAT	TTTCTAATCG	ATATTCATAA	AAATGTCGTG	AAAAGTATG
ATTTCACACG	GAATATGGTT	GAGATCTTGA	AGAAAGTCAA	AGCCCAAGGA	CTCAAGCAAC	CAGTCACTCT	CGCGATTCAA
CGATCTGATT	ATATGTGTCA	TAAGGATCAA	TATTCAGCGG	AATATGGACT	GAAACAAATT	GAAATAAACA	ATATCGCCTC
GTCAATGGGA	GCACATGCTC	TACGGCTCAC	CGAATGGCAT	ATCAGAGTTC	TTAAAGCGTT	GAACATTTC	GATGACGTCA
TTCAAAGAGC	AATTCCAGAA	AACAAGCCAA	TTCCAATGAT	CGCTGAAGCT	TTATTCAAGG	CCTGGTCCCA	CTTTTCGAAC
CCAGCAGCTG	TGGTTCCTGT	CGTTGTAGAA	AACGTCAATC	AAAATCAGAT	TGATCAACGC	CACGTGGAAT	ATGAACTTGA
AAAGTTAGGA	GTACCGATGA	CATGTATTAT	TAGAAGAAAT	TTAACACAAT	GCTATGAACA	ATTATCATTT	AATGATAGAA
GCGATTTGAT	GATTGATGGG	CGTCAAGTAG	CAATTGTTTA	CTTCAGAGCA	GGATACTCAC	CTGATCATA	TCCATCTACA
AAAGAATGGG	AAGCACGTGA	GCGTATGGAA	CTTTCCACCG	CTATCAAAAC	TCCATGGATC	GGGCTACAGG	TGGCAAATAC
TAAGAAGACC	CAGCAGGTTT	TTTCTGAAGA	TGGAGTACTC	GAAAGATTCA	TCGGAAAACC	ACGAGAAGCT	CGCGATATT
GAGCTTCATT	CGCAGGAATG	TGGGCTTTGG	AGAACACTGA	TGAAGTGACT	ATGAAAGTCG	TGGCTGGAGC	TCAAAAACAT
CCAGAAGCGT	TTGTTCTGAA	GCCACAACT	GAAGTGAGG	CCGCATTGCA	CACCGGTGAT	GAGATGGTTC	AAATGCTCCG
AGAAGTTCCG	GAAGAAGAGC	GTGGAGCTTT	CATTTTGATG	GAGAAACTGA	AACCGATGAT	TATTGAAAAC	TACCTGGTTC
TTGCAAAGAA	GCCGATCACA	TTTGCTAAGG	CTGTTAGTGA	ACTTGGAGTG	TATGGTTATG	CATTTGGAAG	GAAGGATGCA
CCTGAGCTTA	AGACTGCTGG	GCATTGCTC	CGAACGAAAC	CGGAATCCAC	AGCTATGGGT	GGAGTAGCCG	CCGGACATGC
TGTTGTCGAC	ACCCCATTC	TCTACGAATT	TATTTGATtt	cgaacataat	cagaaaactc	aacaaaaatg	ctgtgatatg
aaaccatttg	ctatttagat	ctttttgtgt	ttgtaaattt	aatcattgta	atttattgaa	tgt	

FIG. 8



The F39B2.3 gene is located on chromosome I. Regulatory sequences can be found e.g., in the region between 14 797 521 and 14 798 310. An exemplary sequence of this region is as follows:

CATTTTGAAAGTGCCCAAGTTGCTGGAACGCTGAAAATTGAAATTATTAACAAAGAAATTTGCTTTAAAA  
TCCGAAAAATCAAGAAAAAATCGATAATTTTCGTGCGACAATCCGCCTGCTAGCACGGCTTGACGCTCGTT  
TGCCGCGCGCTCATTCGATTTGTGTGAGTGCCCAGTGAGCGCGTTTGCTAAGGCTAACTGTGTAGTCCT  
CTCGACAAGATCTGTGAACATTGAAATGAAACACTTGGGTTCAATAAAATCACAAGAAATGATGACAA  
-518  
TTTGTGTTGCGACCGAAAAAAATTATAAAAATTGAATATTGGTTATCATCGTTTCAATCTTTGTTTTGT  
-469  
ATTAAAGGCACAGCTGCTAAAAATTGTTTTTTTTTTTCAATTTTGCTAAAAGAAAATCAATTTTCTGAT  
TTTTTGTGAGTTCCCGTGCAAATCAATGTCCTAGCTTTTTTAAATTTGTTTTTTGTTATGTAATTCAT  
CAAATTTTGTGCAATTTTCTAGAGATTTTCTGCTAAAACACTAAAAATAGTCTAAAAGTCGATAATTTGAT  
AAACATTTACTCAAACCTTTTACGGAAAAATGAAACAAAAGTTGCAAAAATATAGTAATTTTCGCAATTTT  
CTGAACGCGTACTTAAAGGTACACGGTTTGATTTCGGATTGGTCCCGCCACAAAGTGTACCATAACATTT  
TTCTCGCTGCGAGACCCATCCGAATAAATCCGTGCGCCTAATCAGTGCGAGTACGCATTTTCATATTACTG  
ATAAGTGCCATTTTGAACAATG

FIG. 9

Matter No.: 10861-032P01  
 Applicant(s):  
 SKN-1 GENE AND PROTEIN

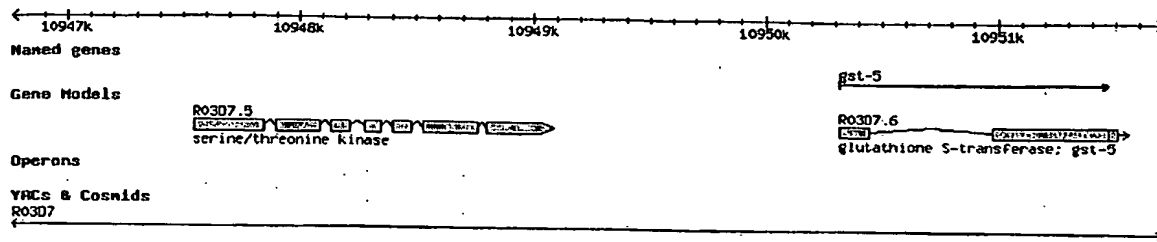
Page 10 of 31

Exemplary F39B2.3 sequences:

Amino Acid Sequence							
MSKSICKSSM	RAAVVRRFGA	PDVIEAVESD	MPRLEKNQVL	VRNYAAGVNP	VDTYIRAGQY	GKLPNLPYVP	GKDGAGFVEL
VGESVKNVKV	GDRVWYGSEA	DSTAEYVAVN	RPFELPEGVS	FEEGASLGVP	YLTAYRALFH	LAGAKTGDVI	LVHGASGGVG
SALMQLAAWR	NIEAVGTAGS	ADGIRFVKSL	GARNVYNHSD	KQYVSKMKND	YPGGFNHIFE	MAAHTNLNTD	LGLLAPRGRV
AVIGNRAETT	INARQLMVTE	GAVYGVALGM	SSEAEILLDFG	INIVSFLKET	EFRPLINKLY	RLEQLGLAHE	EIMNNKGAKG
NLVVQIEH							

Spliced mRNA							
ATGAGCAAAT	CGATTTGCAA	ATCAAGCATG	CGCGCAGCTG	TAGTCCGACG	ATTCCGAGCA	CCTGATGTCA	TAGAAGCCGT
CGAGAGTGAT	ATGCCCAGGC	TTCGAAAAAA	CCAGGTTCTC	GTTCCGAATT	ACGCTGCCCG	TGTCAATCCA	GTTGACACAT
ATATTCGTGC	TGGTCAGTAT	GGAAAACTAC	CAAATCTTCC	ATATGTACCA	GGAAAAGATG	GAGCCGGATT	CGTCGAACTT
GTGGGAGAAA	GCGTTAAAAA	TGTGAAAGTC	GGCGATCGAG	TCTGGTATGG	ATCAGAAGCG	GACAGTACAG	CAGAGTATGT
TGCGGTGAAT	CGACCATTCG	AGTTGCCGGA	AGGAGTTTCG	TTTGAGGAAG	GAGCTTCTCT	CGGAGTGCCT	TATCTTACCG
CTTATCGTGC	ATTGTTTCAT	CTTGCTGGTG	CAAAGACTGG	CGACGTTATA	CTTGATACAG	GAGCATCTGG	TGGAGTGGGA
AGTGCACTGA	TGCAGCTGGC	TGCCTGGAGG	AACATTGAAG	CTGTTGGCAC	TGCTGGATCT	GCTGATGGGA	TCCGGTTCGT
GAAGAGTCTT	GGTGCACGGA	ATGCTATAAA	TCATTCCGAT	AAGCAATATG	TGTCGAAAAT	GAAAAATGAT	TATCCAGGAG
GCTTCAACCA	CATTTTCGAA	ATGGCTGCTC	ACACAAATCT	GAACACGGAC	CTCGGATTGC	TGGCTCCACG	TGGTAGAGTT
GCAGTAATTG	GAAATCGCGC	CGAGACCACG	ATCAACGCAA	GACAACCTAT	GGTTACAGAA	GGAGCTGTTT	ACGGTGTAGC
ATTGGGAATG	TCTTCCGAGG	CTGAGCTCTT	GGACTTTGGC	ATCAACATTG	TCTCATTCTT	GAAGGAAACC	GAGTTTCGTC
CACTTATAAA	CAAATTGTAT	CGTCTCGAGC	AATTAGGACT	GGCTCATGAG	GAAATTATGA	ACAACAAGGG	AGCGAAAGGA
AATCTTGTAG	TGCAAATCGA	ACATTAAATC	attatttttaa	cacgccatth	aaaggaa		

FIG. 10



The R03D7.6 gene is located on chromosome II. Regulatory sequences can be found e.g., in the region between 10949088 and 10950317. An exemplary sequence of this region is as follows :

AGAACTTTTCGAGAAGTCTACCGTTGTAGTTTTTCGAAATAGTAATTTATTTAGTGACGTTTATAAAGGTTTACATGATTT  
GGTTTGGAAATTTTGTAGGAGTTTATTCATAAAACAAAGTAACCATGGACATTCCAGAAGTCTATAGTACACGCGATCC  
TACCGTACCCTTCAGTATTTCTATCAGATTGATAGCTTTCGGTAGTCAGGTACAGCCTAAAAAATTCCTGCTTGCCTTTT  
TGCTTACATGTCTGCCTACCTTCAGTCATAATGCCTACATAATG

- 947

ATTTTTTCCAATTGAACTTGCAGACAGAAATTCAAATGGCAAAAAGAAACAAACACCGAAACATTAATCA

CATTTCTTTTCATATCAGTTTTCTTCTGTCAAAGCACATTTCTGGAGTCTGTGTGTATTTTTTGTGTC'TTTATGTGATCGG  
TGTTGTGAAATTTGTAGTTGATGTTGATAACATACTTTTTTTTGAACAAAAAGTGATTGATTAGGCTTGAATTCAGAGA  
TATGTTTCGTGATAC'TTTGCGATTCTCGAGCCAAAAACACGGTATCCGGTCTCGACACGACAAC'TTTTCGCAAAATACAA  
GCTGATGTGCGCCTTGAAAGAGTACTGTAATTTCAACCTTTCGTTGTGTGCGGAATTTTCATAGTTTCTCGTTCAAAATAT  
ATGTATTTATTAACAAAAAATAAAACAAAAACAATTGAGAACACATAAATGTGAAAAATCAATGAGACCACAGCAAAA  
AATTTTGTATCTACAGTACTCTTTAAAGGCGCACATCCGTTCTTATTTTCAGCAAAAAATGTCGCTTCGAGACCGGGTACC  
GTATTTTTTTTTGTGCAAAACTTTAGGTCTAGGTAATATTAATAAAAAAATTCACAAAACTAGAATCTAGAGCTTTCCAT  
TAAATTTTTTGATGACATTTGAAATTCATGATGATTTTTTTTCCAACAATTCGAAATATCCCTCTTTTCACCTGGTCC

-302

-282

ACTGAATTCCTTTCCGAAAGACCACCACAATTTTCAGGGCTCCGCCCATTTTCGTGGTTTGTAGCCTTCCCGACCCTACGT  
TTTTGATGACAATTGTGAGAGAAGTGAGAGGTTTCAGACACAAAAAGCGACGTGGTCGAATGA

-149

GTATAAATAGAGAGTGAAGTTTCCAATTTCCCTCACAATTGTTTGTGTTGCAATCCACTTTCCAAAAAACACAACCTTCAA  
TCAAAAATCATTATGGTT

FIG. 11

Matter No.: 10861-032P01  
Applicant(s):  
SKN-1 GENE AND PROTEIN

Page 12 of 31

Exemplary R03D7.6 (gst-5) sequences:

### Amino Acid Sequence

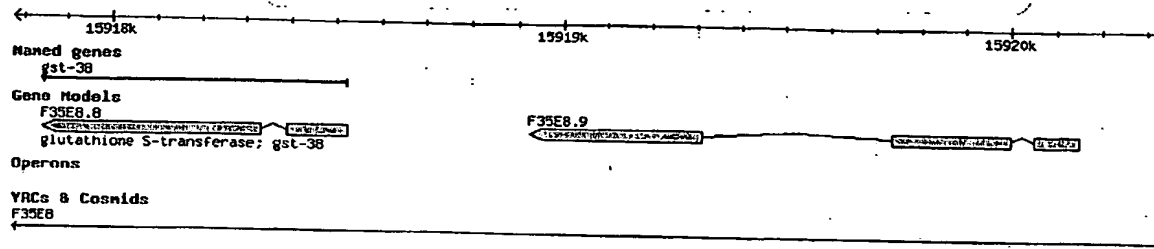
MVSYKLTYFN GRGAGEVSRQ IFAYAGQOYE DNRVTQEQWP ALKETCAAPF QLPFLEVDG KKLAQSHAIA RFLAREFKLN  
GKTAWEEAQV NSLADQYKDY SSEARPYFYA VMGFPGDVE TLKDIPLPA FEKFGFLVN FLKASGSGFL VGDSLWIDL  
AIAQHSADLI AKGGDFSKFP ELKAHAEKIQ AIPQIKKWE TRPVTPF

### Spliced mRNA

ATGGTTTCCT ACAAGTTGAC CTA CTCTCAAT GGACGTGGCG CTGGAGAAGT GTCTCGTCAG ATTTTCGCCT ATGCCGGACA  
ACAATACGAG GATAATAGAG TCATCAGGA ACAATGGCCA GCATTGAAAG AAACCTGCGC TGCTCCATTC GGACAACTTC  
CATTCTCGA AGTCGACGGT AAGAAGCTTG CTCAATCCCA CGCGATTGCT CGTTTCTTGG CTCGTGAGTT CAAGCTCAAC  
GGAAAACCG CCTGGGAAGA GGCTCAAGTG AACTCTCTTG CCGATCAATA CAAGGATTAT TCAAGTGAGG CTCGTCCATA  
TTTCTACGCT GTCATGGGAT TCGGTCCAGG AGACGTTGAA ACTTTGAAGA AAGACATCTT CCTTCCAGCA TTTGAAAAGT  
TCTACGGATT CTTGGTCAAC TTCITGAAGG CTTCCGGGATC CGGATTCCTT GTCGGAGACT CTTTGACCTG GATTGACTTG  
GCTATTGCCC AACATTCAAGC TGATTTGATT GCCAAGGGAG GTGATTTTCA CAAGTTCCCA GAGCTCAAGG CTCATGCCGA  
GAAGATCCAG GCGATTCCAC AAATCAAGAA ATGGATCGAG ACCCGTCCAG TCACACCATT CTAAatagct gtataaaatc  
tgcaataaaa tattttttttt tttt

FIG. 12





The F35E8.8 gene is located on chromosome V. Regulatory sequences can be found e.g., in the region between 15 917 841 and 15 918 925. An exemplary sequence from in or around this region is as follows:

TCTCATTCTCTTCAAGACATAACACAACGGGCTGACGACCATATCATCAACGACGATTTTTTAGGAACTG  
TACTTTATCTGTGTCTGACCAACACGTGTGAATGAAGTTTCAACTGGAAAATTTGTTTGAAACACTGCAA  
AGAATTTGGAATTTTGATGATAATTTTAAATGCCATTATCAGTTTTAATACGCCACTCTAGTCTTTGATT  
-240

CTTTGCACACACACACACACACACACACACACACACTCACAAACACGCCTGAAATTTGCAATATG  
CTGATTTAACGAGAAAACATTTGATGACAATAAACTTGGCGTATTAATATAAAAGGGAATTCATTCA  
-94

GATTCTCAACGGTTTATTTTCTGTCACAACCTCTTCCTAATATTCACCATGGTTT

FIG. 13

Exemplary F35E8.8 (gst-38) sequences:

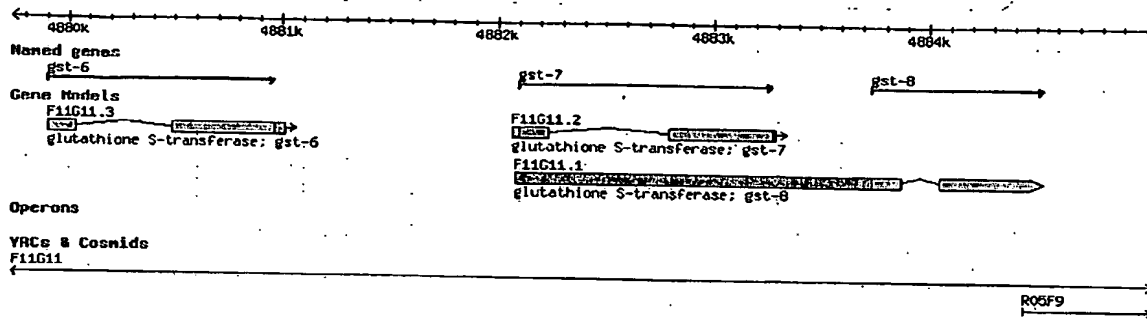
### Amino Acid Sequence

MVSYKLTIFYD GRGAGELCRQ IFAAAEQKYE DNRLTDEEWE KFKAAAGKTPY NQLPMLEVDG KPLAQSHAMA RYLAREFGFN  
 GKSRWEEAQV NSLADQYKDY YAEARPYLAV KLGYTEGDAE ALYTSVYLPV FKKHYGFFVN ALKASGSGFL VGNSLTFIDL  
 LVAQHSADLL GREKSDLFND VPENKAHSEK VQSIPQIKKW IETRPASDW

### Spliced mRNA

ATGGTTTCCT ACAAGCTTAC CTAATTTCGAT GGACGCGGAG CCGGAGAGCT CTGCCGTCAA ATCTTTGCTG CCGCCGAGCA  
 GAAATATGAA GATAACAGAC TTACCGATGA GGAGTGGGAG AAGTTCAAAG CGGCCGGAAG AACCCCATAC AACCAGCTTC  
 CAATGCTCGA GGTAGATGGC AAACCACTCG CTCAGTCCCA CGCGATGGCT CGTTATCTTG CTCGGGAATT CCGGTTCAAC  
 GGAAAGAGCA GATGGGAAGA AGCTCAAGTC AACTCCTTGG CCGACCAGTA CAAAGACTAT TACGCGGAGG CTCGTCCATA  
 CCTCGCTGTG AAGCTTGTT ACACAGAAGG AGACGCGGAG GCTCTTACA CAAGCGTCTA TCTTCCAGTT TTCAAGAAAC  
 ACTATGGATT CTTTGTCAAT GCTTTGAAGG CCAGCGGGTC AGGATTCTTG GTTGGAATTT CCTTGACTTT TATTGATTTG  
 CTTGTTGCTC AGCAATCAGC TGATTTGCTG GGACGTGAAA AGTCGGATCT TTTCAATGAT GTCCAGAGA TGAAGGCACA  
 TTCCGAAAAA GTTCAGTCAA TTCCTCAGAT CAAGAAATCG ATTGAGACTC GTCCAGCGAG TGACTGGTAA

FIG. 14



The F11G11.2 gene is located on chromosome I. Regulatory sequences can be found e.g., in the region between 4 880 968 and 4 882 068. An exemplary sequence in or around this region is as follows:

AATTGAAATGAGTTTGCAATTTTGTATTATTTTTTAATTCATATTTCAAGAAGCAATTTTTTGCTAATTTGTTTTAATGG  
AAATCGATGTTTCTAAAATATCTTGAATGAATTGTTCTTTTAAAAATTTTATGGTAAAGTTTTCAGCAGGATGTTTCTAT  
AGAAGCTTTTTGCATTGCAAGAGTGTTGAAATATACAGGATATTTACAAAAGCCTGGGAAGTAGGCATGCTTTTAGGTAC  
AAATCAGACCTACACCGCTTCTTTGTGGTTTACCATCATAGCTAAACTTTCCGAACATTCCTGGTGAGACACAATG  
TTCAAAGCACAAAACCAATCACGTCATAATGTTAATTTGACTTTTATTGTCAAAAATTACAAAAGCGTCGTTTTCTGGAA  
CATGAACATAATAAGAATTTTCAAATTTCCGGTGGGCACAATAAATATGTAATCTTTTATTTATTTTGGAGGATAGTCTT  
TTCAAAGGCAGGTGTATAACCTCAAAGAAAGCACGTTTGTGTTTCAAAGTGAGACTTAAATTTATTTCAAAGACAAAT  
CCATAGGAAATCATTGTTTCATCAGGCACCTTCCAGAAATTAGGCTGTAGGCAGGCACGTAGGCTGCGGTAAATGCCTAC  
GCCTCTTTTGCGCGAGATTATGAAATTGTGTTGTACTGTGCGGAAAAATTTAGAAACAAAAAAATATTTTGTGACT  
TTTTGTGTCAGTTATAGTAGTTTCTTATCATGGTATCTTCAATAATAATGGCAAGCGTAAC  
-391

AAGATGATTGATGCCATGGGTTTATATTTGTGAGTAGTCACAAATTGTGACACAACATTCCTTCGAAAGATCTGGAAAA  
GTCACAAAACCTTGCATATATTTTTTCAACCAATATTATTTTGACCTACTCTGTTTCATCGTAACATTGCAACAACAAAA  
AACGATGACTACACTTTATGATTTCTAGTCAACAACGTGCGCGCAATGTGTAGAGCAAATGATGACAACTACAGAATAT  
GGTGAGTGGAGAGACGACAGACATTTGAGAAATGGGTATAAATA  
-133

GAGACGGCCGGCATTCAAGTGTCAACCTTCTCATCGACCACTCGATTTCTTGCTTGTTATTTCAACAATG

FIG. 15

Matter No.: 10861-032P01

Page 16 of 31

Applicant(s):

SKN-1 GENE AND PROTEIN

Exemplary F11G11.2 (gst-7) sequences:

**Amino Acid Sequence**

MVHYKVSYPF IRGAGEIARQ ILAYAGQDPE DNRIPKEEWP AVKPSTPFGQ LPLLEVVDGKV LAQSHAIARY LARQFGINGK  
 CAWEEAQVNS VADQFKDYLN EVRPYFMVKM GFAEGDLAL AKDVFLPGFK KHYGFFANFL KSAGSGYLVG DSLTFVDLLV  
 AQHTADLLAA NAALLDEFPQ FKAHQEKVHS NANIKKWLET RPYTPF

**Spliced mRNA**

cgaccactcg atttcttgc tggttatttc aacaATGGTC CACTACAAGG TATCGTACTT CCCAATTTCGT GGAGCTGGAG  
 AGATTGCTCG TCAGATCTTG GCCTACGCTG GACAAGACTT CGAGGACAAC AGAATCCCAA AGGAGGAATG GCCAGCTGTC  
 AAGCCAAGCA CTCCATTCCG ACAGCTTCCA CTCCTTGAAG TTGACGGAAA GGTTCTTGCC CAATCTCATG CTATCGCCCCG  
 TTAATTGGCT CGTCAGTTCG GAATCAATGG AAAGTGTGCA TGGGAGGAGG CTCAAGTCAA CTCGGTTGCT GATCAATTCA  
 AGGATTACCT CAACGAAGTT CGTCCATACT TCATGGTGAA GATGGGATTT GCTGAAGGAG ATCTCGATGC TCTTGCCAAG  
 GACGTCTTCC TTCCAGGATT CAAGAAGCAC TATGGATTCT TTGCTAACTT CCTCAAGTCG GCTGGATCCG GATACTTGGT  
 TGGAGACTCT TTGACCTTGG TCGACTTGCT CGTCGCTCAG CACACTGCTG ATCTTCTGGC TGCCAACGCA GCTCTTCTCG  
 ATGAATTCCC ACAATTCAAG GCTCATCAGG AAAAGGTTCA CTCGAATGCC AACATCAAGA AGTGGTTGGA GACTCGTCCA  
 GTTACTCCAT TCTAAatgat ttcca

FIG. 16

Matter No.: 10861-032P01  
Applicant(s):  
SKN-1 GENE AND PROTEIN

Page 17 of 31

The K08F4.7 gene is located on chromosome IV. Regulatory sequences can be found e.g., in the region between about 10141800 and 10142217. An exemplary sequence of this region is as follows:

ATTATCCAAAAAGATTAGAAAGTTGGCAAACCTTGGGCAAGAATTTCCAGAGATTGCACTAAAGTTGTAGCCAAGTTTGAT  
CCAACTTTATCCAATCTTTTACTAAAATTATCCTTAAGACTATTTAAATTTTAGATAGAGAATTGGCGAGAGTTAGATCC  
CACTTGGATATGACTTATAGTTAGCCTAACCTGAAGCTATTGCTTGCTTGATCATTGGTTTATCGCTTTGCTACTTGGA  
TAACCAGCTCCAATAGTTGTTATTTTGTCTTTGTCATCATTTT

- 157

TCCACGATTTACACTCTCAAGTGAAACCAACTGTTCTTTGATGCCAGACGATGACATTACACTTGATAAGA

- 83

AAATATATATAAACTGGAATTAAAAACAATTGATACATCGATTCAATTACTGAATTCTAATTATG

**FIG. 17**

Exemplary K08F4.7 (gst-4) sequences:

**Amino Acid Sequence**

MPNYKLLYFD ARALAEPIRI MFAMLNVPYE DYRVSVEEWS KLKPTTPFGQ LPILQVDGEQ FGQSMSITRY LARKFGLAGK  
TAESEAYADS IVDQYRDFIF FFRQFTSSVF YGSDADHINK VRFEVVEPAR DDFLAIINKF LAKSKSGFLV GDSL TWADIV  
IADNLTSLK NGFLDFNKEK KLEEFYNKIH SIPEIKNYVA TRKDSIV

**Spliced mRNA**

ATGCCAACT ATAAGCTATT GTATTTTGAT GCTCGTGCTC TTGCTGAGCC AATCCGTATC ATGTTTGCAA TGCTCAATGT  
GCCTTACGAG GATTATAGAG TTTCAGTGGA AGAATGGTCA AAGCTGAAGC CAACGACTCC ATTTGGCCAG CTTCCTATT  
TACAAGTCGA TGGAGAACAA TTCGGTCAGT CAATGTCTAT CACAAGATAC TTGGCAAGAA AATTGGACT CGCTGGAAAA  
ACTGCAGAGG AAGAAGCTTA CGCTGATTCA ATTGTAGATC AATACAGAGA TTTTCATATC TTTTCCGTC AATTCCTTC  
TTCCGTTTTT TATGGAAGTG ACGCTGATCA TATTAACAAA GTACGTTTTG AAGTTGTTGA ACCAGCCCGT GATGATTCT  
TGGCAATAAT CAATAAGTTC CTGGCCAAGA GTAAATCAGG ATTCCTCGTT GGAGACTCAT TGACTTGGGC TGATATTGTG  
ATTGCTGACA ATTTGACAAG TCTCCTGAAG AATGGATTCT TAGATTTCAA CAAAGAAAAG AAGTTGGAAG AGTTCTATAA  
CAAGATTCAT TCAATTCCAG AAATTAAGAA TTACGTGGCA ACAAGAAAGG ATAGTATTGT TTAAaatcga attatttaag  
tctgaattat gtatgtagta aaataatatc gttcctatca cgtctcccag agagcgtaat aaattattat tatgtg

FIG. 18

Matter No.: 10861-032P01  
Applicant(s):  
SKN-1 GENE AND PROTEIN

Page 19 of 31

The *sod-1* gene is located on chromosome II. Regulatory sequences can be found e.g., in the region between about 6 973 806 and about 6 974 406. An exemplary sequence of this region is as follows:

ATTCCGCAACCCCGTCAAATTTAAGAAGAGAAAGAAAAAACAACCGTGTTCACCTGTAAGGTAGT  
TTTTTTTTGTTGCCTTCGGCGTTTTGATTCACATGAAAGTTTCTACGGAAAAACTTTCATTGCATAACGA  
TCTTCATATCTTGTCTTGAAACGAAATTTCCAACATGAAAGAAACCCGACGCTATTTATTCTCGCAA  
CACAAAAATTCACATTTAAATAACCGCGTTTTTCTCGAACAGCATATTTGACGCGCATGCTCGTCAA  
GTTTGATGCGTGCACACTATTTTGCTGTTGTTTTTTCTTTTTCTCTAAATTTCTTTACGCTTTCGTA  
GTTTCTATAGAAACGATTCTCCACTCCCGGTTTTCTCCGATTCTCAAAATTAATTAAAAATTTAGTTATT  
AAAAATCCTTTTTCTTGAAATAATCGTTCAATTTTCGAGTTTTCAAGAGTGGAGACGTTGAATTTGTGAGC  
CGCTTATTTTTCTGTGTTTTTGTGTTTTTAATCAGTGTCATAATCATACTTTCCATTGTTTCT

- 64

TTATTATTCAAAGTTGTAGATTCAGTATTTTAGATCGGTGATG

FIG. 19

Matter No.: 10861-032P01  
 Applicant(s):  
 SKN-1 GENE AND PROTEIN

Page 20 of 31

Exemplary sod-1 sequences:

### Amino Acid Sequence

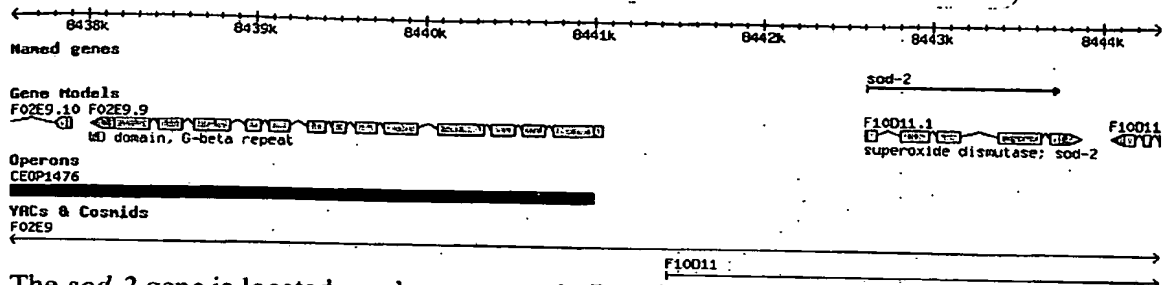
MFMNLLTQVS NAIFPQVEAA QKMSNRAVAV LRGETVTGTI WITQKSENDQ AVIEGEIKGL TPGLHGFHVH QYGDSTNGCI  
 SAGPHFNPFQ KTHGGPKSEI RHVGD LGNVE AGADGVAKIK LTDTLVTLYG PNTVVGRSMV VHAGQDDLGE GVGDKAESK  
 KTG NAGARAA CGVIALAAPQ

### Spliced mRNA

tttagatcgg tgATGTTTAT GAATCTTCTC ACTCAGGTCT CCAACGCGAT TTTCCGCAG GTCGAAGCCG CTCAAAAAAT  
 GTCGAACCGT GCTGTCGCTG TTCTTCGTGG AGAAACTGTT ACCGGTACTA TCTGGATCAC ACAGAAGTCC GAAAATGACC  
 AGGCAGTTAT TGAAGGAGAA ATCAAGGGAC TTAATCCCGG TCTTCATGGA TTCCACGTTT ACCAATATGG TGATTCCACC  
 AACGGATGCA TTTCTGCCCG TCCACACTTC AATCCATTG GAAAGACTCA TGGTGGACCA AAATCCGAGA TCCGTCACGT  
 AGGCGATCTA GGAAATGTGG AAGCTGGAGC CGATGGAGTG GCAAAAATCA AGCTCACCAG CACGCTCGTC ACGCTTTACG  
 GTCCAAACAC TGTCGTTGGC CGATCTATGG TTGTTTATGC CGGACAAGAC GACCTCGGCG AGGGAGTCGG AGACAAGGCA  
 GAAGAGTCCA AGAAGACTGG AAACGCCGGA GCTCGTGCTG CCTGCGGTGT CATTGCTCTC GCTGCTCCCC AGTGActacc  
 tgaatcgct ctctgaatct ccacacaatt cctactaaag acaatttttc atttcttgct ttgtcgttat attcttaaga  
 atccccgttg ttctactcct actactgtat attttcacat aaaatttttc caaaatttca aataaagggt gtagtttc

FIG. 20





The *sod-2* gene is located on chromosome I. Regulatory sequences can be found e.g., in the region between about 8 441 038 and 8 442 612. An exemplary sequence of this region is as follows:

TGAATAAAAAACGTTGAACCCAACGGACATCAAAGTATCAAAGTAAGTAAGTAAGTAAGTAACCTGAATAAAAAACGTTGCA  
TATAAAAAATCTACTCGAAAATTAAGTGAGAATTGAAGGATTGCTTTCCGAAGAGAAAATGACAATTATAGGGTATACTA  
AAACATCAAAAATGTATATTAGACTACCATAAAATATAAAACATCAGTGCTGCTCTCCAAGCTATTCTGACGGATTGCGAC  
AACGAGCTCGCTGGAGTTGGCATCAGTGTTGGAAGGCAGACACATAAGAAGACTCGAATTTGCGGATGACGTAGTCCTGAC  
ATGTTCCACACCGGGAGAAGTTCAAGAACGACTGGAATTTTGGACCGAATAAGTTCTAATTACGGACTCAAGATCAATC  
AGTCAAAGACTGTTCTTCTGAAGAACAAGTTTGGCCGGAGCCAAGACGTCCTTTTCAACGGATCCCCCATCATTCCCGTG  
CCTGGTTGCCGCTATCTGGGTCGCTGGATCGACATTTCTGGCTCAATTGACGAAGAGATCTCGAGGAGAATAAGAGCAGG  
TTGGGGTGCTCTGGTTGGAATCAAAGAAGTCTTGAGAATCATGCCAAACAAGAAAGAATCATCCTCTTCAAGCAAAAT  
- 980 - 959

GTGCTACCAGCTCTCCTGTATGCTAGTGAAACTTGGACTTGTAATGCTGGATCCACGTTGAGACTCAAAAGAACTGTCAC  
CGGTCTCATCGACGCTGCAGAAATTCGAGGCTGGAACCTTCAACTTGGAAACGTTACCTCCTTGCAAAACAATCAAGATTTG  
CAGGACACATTCTACGGAGAGATCCAAACCGATGGACAAAATCTGCACGGAATGGGACCCGAGCCACAACAAAATTTGG  
AAACGTGCCGTTGGAGGACAGAAGAAGAGATCGGCTAAGGACATCGACGAAGAATACGCAAAATTCACCACAATTCCGC  
CATGTCCGGACAAGTCGTTGTTGGGAGAAGAAGACTAGGAATGCTCACTCCGAAGGCTCCATGGCTGTCCATCGCACGAA  
CCGACCGTGAAAAATGGAAAGAGTTTGTCCGAGTTGCCTCGCAACTTGAACCCAACGGACATCAAAGTATCAAAGTAAG  
TAAGTAAGTAAGTAACCTGAATAAAAAACGTTGCAATTAAAAATCTACTCGAAAATTAAGTGAGAATTGAAGGATTGCTT  
TCCGAAGAGAAAAATGACAATTATAGGGTATACTAAAACATCAAAAATGTATATTAGACTACCATAAATATTACGATAAT  
- 363

TTAAAAATTACTAGAAACACGCAATTCGGCTCAAAAAGCAACAATTTAGACTGAAAACGAGCTAAAAGAATATTATTCAA  
AAACCACTTTGCTCGGTAAATCTGGTGTATCATGTTCCGCAACACTGTCTTTTGTGTTTTCG  
- 191

TACTTTGTTTACGCGCATTGCAATTTTCAAGTGTTCGCGCTTTTGTGTTTACTTTTTTATTTTTTATCCAAAAATCGTATTTT  
CAGCTTGATATGTTTCTGCGAATTGTAAAAATTTATATTGACTATTGAATATTTTAATTATTTGCAGCCGAAAATG

FIG. 21

Matter No.: 10861-032P01  
Applicant(s):  
SKN-1 GENE AND PROTEIN

Page 22 of 31

Exemplary sod-2 sequences:

### Amino Acid Sequence

MLQNTVRCVS KLVQPIITGVA AVRSKHSLEP LPYDYADLEP VISHEIMQLH HQKHATYVN NLNQIEEKLH EAVSKGNVKE  
AIALOPALKE NGGGHINHSI FWTNLAKDGG EPSAELLTAI KSDFGSLDNL QKQLSASTVA VQSGSGWGLG YCPKGKILKV  
ATCANQDPLE ATTGLVPLFG IDVWEHAYYL QYKNVRPDYV NAIWKIANWK NVSERFAKAQ Q

### Spliced mRNA

tttgcagccg aaaATGCTTC AAAACACCGT TCGCTGTGTC TCAAAGCTTG TTCAACCGAT CACAGGAGTC GCTGCTGTTC  
GCTCGAAGCA CTCGCTGCCA GATTACCAT ACGACTATGC TGATTGGAG CCTGTAATCA GTCACGAGAT TATGCAACTT  
CATCATCAAA AGCATCATGC CACTTATGTG AACAATCTCA ACCAAATTGA GGAAAAGCTT CACGAGGCGG TCTCCAAAGG  
AAACGTCAAA GAAGCTATCG CTCTTCAGCC AGCTCTCAAG TTCAATGGAG GAGGACATAT CAACCACTCC ATCTTCTGGA  
CTAATTGGC AAAGGACGGA GGAGAACCAT CGGCGGAGTT GCTCACCAGCA ATTAAGAGCG ACTTCGGATC TCTGGATAAT  
CTTCAAAAAC AGCTTTCGGC ATCAACTGTC GCTGTTCAAG GATCAGGATG GGGATGGTTG GGATACTGTC CAAAGGGAAA  
GATCTTGAAG GTTGCCACAT GTGCCAATCA GGATCCACTT GAGGCAACAA CTGGACTTGT TCCACTGTTT GGAATTGACG  
TCTGGGAGCA CGCTTACTAC TTGCAGTACA AGAATGTTTC ACCAGATTAT GTCAATGCTA TTTGGAAGAT CGCCAACCTG  
AAGAACGTCA GCGAGCGTTT TGCAAAGGCA CAGCAATAAa tgagctgaat cacaagaatt aatcgtcaaa tgtagcagta  
gaagttgact cccattgttt tgtaactatt tttgtttcct aattatttcg aaatgtaaat tttcaaacct tttcaaatga  
aaagttttca ccg

FIG. 22

GCACCGCGACTGGGAGTATAAGAATCGCCGAAACATCAATAATCAGTTCGGTAGAAGTGAAAATTGAG  
CGTAAATATGATCATTTTTTCGATGCACCATATTTGACGCGCAATACTTCTACAAGCCGCTGTGTACTGC

TCGTGGACAACCTTTGGATTATTTTTTGTGTTTTTAAAAATTCAAATAGTCAATATATTGCTTATTTATAGCG  
CGCCTTTTTGACAGTAAGTTTGTCAAATTTGCGCGTAAGTTATGGTGTGTTGCACATATGCACCATACAGC  
AACACCCCGCGSCCGGTAGTGGTACATCCATGCAAATGCGCTCTACTGATAATTTGAGTTTAACCAGG  
TTTAGGCGCAAGATAAGAAAAAAGCTTTGGACCAAAAAATTTAGAGTTTATTTTTTTTCGGACATTTTTTA  
TATACATCACAAAAATATGGGCCACTCGTTTTTGATAAAAAACGACAAGCCCAAAAGTTTCAGGTATACGG  
TAGACAAATTCGGTACAGGTACCACTTTTCCACGTAGTGCCAGGTGTGCCATTACGCTTTTGATCTATGA  
AAAATGCGGGAATTTTTTCGTCCAGAAAAATGTGACGTGACGACGTTCTCAACCATGCGAAATCAGTTGAA  
AACTCTGCGTCTATCTCCCGCATTTTTTGTAGATCTGTAGATTTGTAGATCAATCCATTCCCCGTATAC  
CCTGACCCCAATAATCAATACCTACCTAATTTTTGTCTTTCCCCCTACTTTTTTGCCTGTCCAAAATAAGCG  
AGACTATGCCGTAGTCTGTTGCTGCCAACCAATGTTTCCTTATCAGTGATAACGCTACAATCTTCTTTCTTT  
TTTCTCTGTTTCTCTGTGCTCTCCCAACCCATTTCCGTATTACACCTCGTCGTGGTCATTTTTTTTGTTT  
AGAGTTTTATTTAATTCTAAATTTCTAACTAAAATTTTCCAGA

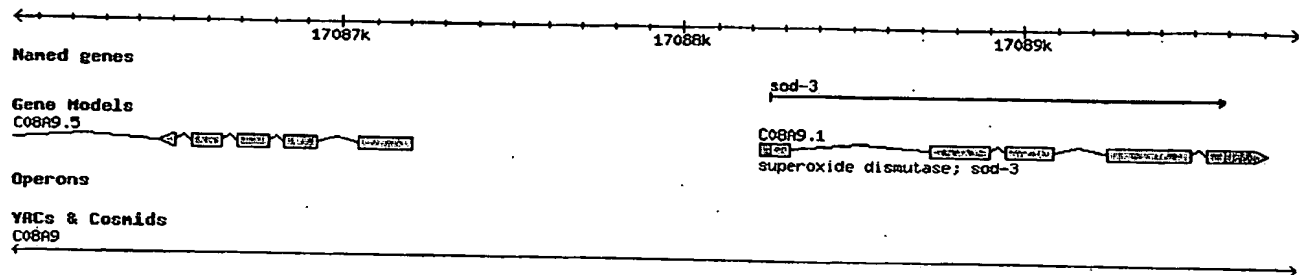
FIG. 23

Exemplary ctl-1 sequences:

Amino Acid Sequence									
MPNDPSDNQL	KTYKETYPKP	QVITTSNGAP	IYSKTAVLTA	GRRGPMLMOD	VVYMDEMAHF	DRERIPERVV	HAKGAGAHGY		
FEVTHDITKY	CKADMFNKVG	KQTPLLVRFS	TVAGESGSAD	TVRDPGRGFL	KPYTEEGNWD	LVGNNTPIFF	IRDAIHFPNF		
IHALKRNPQT	HMRDPNALFD	FWMNRPESIH	QVMFLYSDRG	IPDGFRFMNG	YGAHTFKMVN	KEGNPIYCKF	HFKPAQGSKN		
LDPTDAGKLA	SSDPDYAIRD	LFNAIESRNF	PEWKMFQVM	TFEQAEKWEF	NPFDTVTKWVP	HGDYPLIEVG	KMVLNRRNVKN		
YFAEVEQAAF	CPAHIVPGIE	FSPDKMLQGR	IFSYPDTHYH	RLGPNYIQLP	VNCPYRSRAH	TTORDGAMAY	ESQGDAPNYF		
PNSFRGYRTR	DDVKESTFQT	TGDVDRYETG	DDHNYEQPRO	FWEKVLKEEE	RDLRVGNLAS	DLGGCLEEIQ	NGMVKEFTKV		
HPDFGNALRH	QLCQKKH								

Coding									
CTGAAACCT	ACAAGGAGAC	GTATCCAAAA	CCCCAAGTGA	TCACAACTTC	aaaATGCCAA	ACGATCCATC	GGATAATCAA		
CGTGCTCACC	GCCGGGCGGC	GTGGCCCAAT	GCTCATGCAA	GATGTAGTTT	AAATGGAGCT	CCGATCTACT	CGAAGACCGC		
AACGTATCCC	CGAGCGTGTC	GTTCATGCCA	AGGGAGCCGG	AGCCCATGGA	ATATGGATGA	GATGGCTCAT	TTCGATCGTG		
TACTGTAAGG	CCGATATGTT	CAACAAGGTC	GGAAAACAGA	CACCACTTCT	TACTTCGAGG	TCACCCATGA	CATCACCAG		
GGGATCCGCT	GATACTGTCC	GCGATCCACG	TGGATTCTCT	CTCAAATTCT	CGTTCGTTTT	TCAACGGTCG	CTGGAGAATC		
GAAATAACAC	TCCGATCTTC	TTCATTCTGT	ACGCAATCCA	CTTTCCGAAT	ATACCGAGGA	GGGTAACCTG	GATCTGGTTG		
ACTCACATGA	GGGATCCGAA	TGCGCTCTTC	GATTTCTGGA	TGAATCGCCC	TTCATTTCATG	CCCTGAAGCG	CAATCCACAG		
CTCGGATCGT	GGAATTCCTG	ATGGATTCCG	TTTTATGAAT	GGATACGGAG	TGAATCCATT	CATCAGGTGA	TGTTCCCTTA		
GAAATCCGAT	TTATTGTAAA	TTCCATTTCA	AGCCTGCTCA	AGGTTCCAAG	CGCATACTTT	CAAGATGGTC	AACAAGGAGG		
GCCTCTTCGG	ATCCAGACTA	TGCGATCCGC	GACCTGTTCA	ATGCCATTGA	AATCTCGATC	CAACTGACGC	TGGAAAGCTC		
CATTCAAGTG	ATGACATTCT	AACAAGCTGA	GAAATGGGAG	TTCAATCCAT	GTCAAGAAAT	TTCCCGGAAT	GGAAGATGTT		
ATTACCCACT	GATCGAGGTC	GGCAAGATGG	TGCTGAACAG	GAATGTGAAG	TTGATGTCAC	TAAAGTTTGG	CCACACGGTG		
TTCTGCCCGG	CCCACATCGT	CCCAGGAATC	GAGTTCCTGC	CAGACAAGAT	AATTATTTCG	CTGAGGTCGA	ACAAGCCGCC		
CACGCATTAC	CATCGCCTTG	GACCAAACCTA	CATTTCAGCTT	CCAGTCAACT	GCTCCAAGGG	CGTATCTTCT	CCTACACGGA		
AACGCGATGG	TGCAATGGCT	TATGAAAGCC	AGGGAGATGC	GCCGAATTAC	GCCCCGTACG	CTCCCGTGCT	CATACCACTC		
CGTGATGATG	TGAAGGAGTC	GACATTTTCT	ACGACTGGAG	ATGTTGATCG	TTCCCGAACA	GTTTCCGCGG	ATACCGTACT		
GCAGCCACGT	CAGTTCTGGG	AGAAAAGTGCT	CAAGGAGGAG	GAGAGAGATC	TTATGAGACT	GGAGACGATC	ACAACCTACGA		
GTGGCTGTTT	GGAGGAAATT	CAAAATGGAA	TGGTCAAAGA	GTTTCACGAA	GGCTCGTTGG	GAATTTGGCT	AGTGATTGCG		
CATCAGCTCT	GCCAGAAGAA	GCATTAAatt			GTTTCATCCG	ATTTCCGAAA	TGCTCTTCGC		

FIG. 24



The *sod-3* gene is located on chromosome X. Exemplary regulatory sequences include:

TATTCGCAGAAAAAGTCGTTGCAAACATTTCGTTTTTATATGTTTTTCTTTGAGAAAGCGTGGTTTCATTT  
 TTGAAAGTGAAAAATATTTGCTTAAACTTCCAAATTTAAATCTGCAGTGATTGAGAGAGGTTGAGAATT  
 ATTTTCAAAAACATTCAATGTTTTCCCTTGGAGTGACTATGCAAATATGAAAATGTTTTCCAAAAATATT  
 TGGATGCCCTGATAAAAAGTAGGTGAAATTTTCGCAGGGGAACATCATATTAAAATGTTGAATTTTTAGAA  
 GAAATGGAAATGTTTGTGCGGTGGTATGCTCGAATATTTGAGATATTATATATTTACTGTTAAATCCGAAA  
 TTTTGTGACAAACGGAAAAAATTTGTGTGAAATACTACATTTTCGATAACACAAAGGTACTTCCATAACA  
 CTTATAAAAACGTTTGGACTATCTTATTTTCAGGAAAAAAAATCCAAGAATAAACATTTTTCAGAATTTG  
 AACTTTCTAATGGCTGATTAATAAAAACAAAGTTATACAACATTTCAAAGCAGTTGCTCAATCTGGCATTT  
 TCTTGTGTTTTTTTTTTGAATATTTTCATCAGCAAGATGTTGATAATTTGTGTTAATCTAATTGTTTTCT  
 ACAATTTTTCAAACCGAAAATTGACCTTTGACTTTGTTTACTTTGTTCTCGTGGGTTAACTGTTCACTGA  
 TTTCTATTGCTGTTGATGAGGTCTTTGATCAAATTTGTATTGTTTTTATACTGCATATTGCTTCAATTCT  
 AAATCATCTAATATATTGTCAAACAACCTCTTGTTTTTTTTTTTCATTCAAACCTCTGCAAAAACGTTCT  
 -287

CTTAACAAAGGTTACACAACAACCTCTCCTCTCCATCTCTTTCTCTCAACAACAATGTGCTGGCCTTGCA  
 TGTTTGCCAGTGCGGGTTGTTTACGCGTTTCAAGATTTTGGTCTCCTATCTAACGTCCCGAAATGCAT  
 TTTTCTCTTTCATTTGTTTTTTTTCTGTTGAGAAAAGTGACCGTTTGTCAAATCTTCTAATTTTCAGTG  
 AATAAAATGCTG

FIG. 25

Exemplary sod-3 sequences:

### Amino Acid Sequence

MLQSTARTAS KLVQPVAGVL AVRSKHTLPD LPFDYADLEP VISHEIMQLH HQKHHTATYVN NLNQIEEKLH EAVSKGNLKE  
 AIALQPALKF NGGHHINHSI FWTNLAKDGG EPSKELMDTI KRDFGSLDNL QKRLSDITIA VQSGSGWGLG YCKKDKILKI  
 ATCANQDPLE GMVPLFGIDV WEHAYYLQYK NVRPDYVHAI WKIANWKNIS ERFANARQ

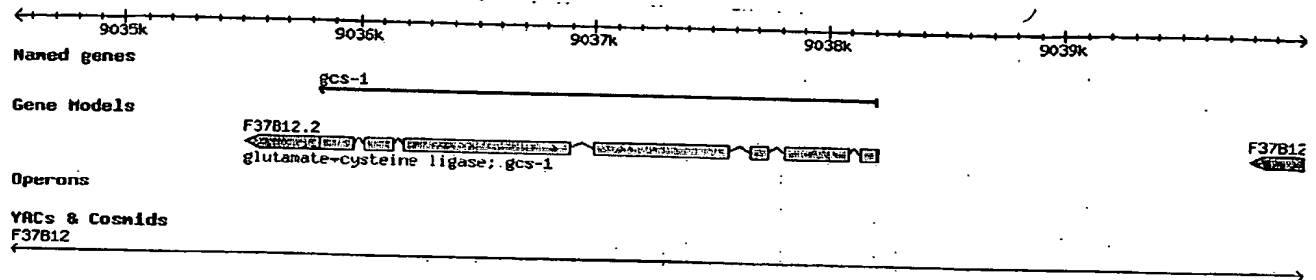
### Spliced mRNA

cgtttgtcaa atctttctaatt tttcagtgaa taaaATGCTG CAATCTACTG CTCGCACTGC TTCAAAGCTT GTTCAACCGG  
 TTGCGGGAGT TCTCGCCGTC CGCTCCAAGC ACACCTCTCCC AGATCTCCA TTCGACTATG CAGATTTGGA ACCTGTAATC  
 AGCCATGAAA TCATGCAGCT TCATCATCAA AAGCATCATG CCACCTACGT GAACAATCTC AATCAGATCG AGGAGAAACT  
 TCACGAGGCT GTTTCGAAAG GGAATCTAAA AGAAGCAATT GCTCTCCAAC CAGCGCTGAA ATTCAATGGT GGTGGACACA  
 TCAATCATTC TATCTTCTGG ACCAACTTGG CTAAGGATGG TGGAGAACCT TCAAAGGAGC TGATGGACAC TATTAAGCGC  
 GACTTCGGTT CCCTGGATAA CTTGCAAAAA CGTCTTTCTG ACATCACTAT TGCCTGTTCAA GGCTCTGGCT GGGGATGGTT  
 GGGATATTGC AAGAAAGACA AAATCTTGAA GATCGCCACC TGTGCAAAACC AGGATCCTTT GGAAGGAATG GTCCCACTTT  
 TTGGAATTGA CGTTTGGGAG CACGCCTACT ACTTGCACTA CAAAAATGTC CGCCCACTAT ATGTCCATGC TATTGGAAG  
 ATTGCCAACT GGAAGAATAT CAGCGAGAGA TTTGCCAATG CTCGACAATA Aaagcaggaa atattggaat tttcggtttt  
 acgaaaatat tgaagataat tcagatgtag tttaaaacgc tgagaatttg tatttttgta attgttttaa taaaagaacg  
 cacagttttt tctta

FIG. 26

Matter No.: 10861-032P01  
 Applicant(s):  
 SKN-1 GENE AND PROTEIN

Page 27 of 31



The *gcs-1* gene is located on chromosome II. An exemplary regulatory sequence is:

TTATCAACCACTAGGTTCCGTCTTAATCGTCCAAATATTGATCCGCTCGCTCGTGTCTTCTCAACTTCTTTATTTGCTGT  
 GTTTTTCTGTTTCTATAGTTCTCCATTTTCCATCTCCTCTTCGCTTGTTGAATGGACTTTATTTTGATAAGTTCATTTTA  
 ATTTTCTAACAATCTCATCACTAGCTCATGATGACAATTGCAAAGAAATTCGTATATAGAGGGGAAAAATGCTGACAA  
 -607

ATATTGAAAAGCCTTCAGGAGAGATGTAGAGACGTAGGAGTAGAGACAGAACATAAATTTGAGAAGCTTGTAGGGAGAAT  
 AGACATAGAGTTACCATGGGAAAAACGCTCGCATTTTCCATTTAACGAGATTTTCTAGATCACAACATTTTGTGATCCGT  
 TGTGCGAAAATCAAGCTTTTATCAAACCTTTATCGTCTGTTTCATTCTTTCTGACAATCTTTATTATCTTATTAAACTTG  
 ACTAATTGTATTGAAAGTATTTTTTTAGATGCGAACGAAGTTCATTTTTTCATGACTTAACATCTCTTAACGTTAGTGAA  
 -316

ATTTTGAATTCCAATTAGGACTACGGTAGGAGTCTGTAGTTGATTTTCCTGAACACTTGTTTTGTAACCTTTCTGAACG  
 GATTTTAATATTTCTAAAATTTTAAATTGCAAATCTGAGTCCTATTAAAAGATGTTTCATCCGTAAAACCAACAAACAAA  
 ATATCACTTTATCATCATGAGATTTAATGTTTCCTTTTGATTTCTGAATTGTTGTACTTTCCTTCAAACGACTTATTGA  
 -121

ACTGATGTAACCTTCCTTCTAATGTTATCATTTGTATTTTTTTGCAGAATG

FIG. 27

Exemplary gcs-1 sequences:

Amino Acid Sequence									
MGLLTGKSPL	TWAETVPHID	YIKKHGIAQF	INLYHRLKSR	HGDQLKWGDE	IEYTIVKFDD	ANKKVRVSCK	AEELNKLQA		
EEQVNAMLGT	ANRFLWRPEF	GSYMIETPG	MPYGGLIACF	NIVEANMKLR	ROVVKKLLKK	DETCLISFP	SLGVPGFTEP		
EVAADRKNDD	AANSVFWPEQ	AVFLGHPREF	NLTKNIGRR	GSKVAINVPI	FKDNTNPSPF	VEDLSALGGP	DDTRDAKPDH		
IYMDHMGFGM	GCCCLQVTFQ	AVNVDEARWL	YDQLTPITPI	LLALSAAPI	FRGKLSNVDS	RWDIISASVD	DRTPEERGLE		
PLKNSKWVID	KSRYDSTDCY	IYPCSVGYND	IPLQYDETIY	KQLIDGNIDE	PLAKHIAHMF	IRDPHQVFRE	RIEQDDEKSS		
EHFETIQSSN	WMNMRFKPPP	PDAPETGWRV	EFRPTEVQLT	DFENAAAYCCF	VVLLTRMMIS	FRITYLMPIS	MVTENMKRAQ		
QKDAVLNQKF	LFRKGLAECK	SAPENLKGSE	KCGPPSQDIE	EMSIDEIING	KKNQFPGLIS	LIRQFLDSAD	VDVDTRCTIS		
QYLNFIKRA	TGEINTLAHW	TRGFVQSHPA	YKHDSVDNDN	IVYDLLKKMD	AINNGEDHCE	KLLGCYRSKT	DHAISAAVRK		
AEEHMIVSSQ	KRAH								

Spliced mRNA									
tttgcagaAT	GGGTCTTTG	ACGAAAGGTA	GTCCGTTGAC	GTGGGCAGAA	ACCGTACCGC	ACATTGATTA	TATCAAGAA		
CACGGAATTG	CTCAATTCAT	CAATCTCTAC	CATCGTCTGA	AATCAAGACA	CGGAGATCAA	TGAAATGGG	GAGATGAGAT		
TGAATACACT	ATTGTAAAT	TTGATGACGC	AAACAAGAAA	GTTCGCGTGT	CGTGCAAAGC	TGAAGAGCTT	CTTAATAAGT		
TACAAGCCGA	AGAGCAGGTG	AATGCGATGC	TTGGAACATGC	CAATCGATTTC	CTTTGGAGAC	CAGAATTCGG	ATCCTACATG		
ATCGAGGGAA	CCCCCGGAAT	GCCTTACGGA	GGTCTCATCG	CTTGCTTCAA	CATTGTCGAG	GCAAACATGA	AATTGCGCAG		
ACAGGTCGTC	AAAAAGTTAT	TAAAGAAGGA	TGAAACATGT	CTATCGATAT	CGTTCCCATC	TCTTGGAGTA	CCTGGATTCA		
CATTCCCGGA	AGTAGCAGCT	GATAGAAAGA	ATGATGATGC	AGCTAATAGC	GTTTTCTGGC	CAGAACAAGC	TGTATTCTTG		
GGCCATCCAC	GTTCGAAGAA	TCTTACCAAA	AATATTAAG	GTGCGAGAGG	AAGTAAAGTA	GCTATCAACG	TCCCGATATT		
CAAGGATACG	AACACCCCCA	GTCCATTTCGT	TGAAGATTTA	TCTGCACTTG	GAGGTCCTGA	TGATACTCGT	GATGCGAAAC		
CTGATCACAT	TTATATGGAT	CATATGGGAT	TCCGAATGGG	GTGCTGTTGT	CTTCAAGTCA	CTTTCCAGGC	TGTGAACGTC		
GATGAAGCCA	GATGGTTGTA	CGATCAGCTG	ACACCGATTA	CACCGATTCT	ACTGGCACTC	TCTGCCGCCA	CACCAATCTT		
CCGTGGAAAA	TTATCCAATG	TGGATTCTAG	ATGGGATATC	ATTAGTGCAA	GTGTCGACGA	TGCTACACCG	CAGGAAAGAG		
GATTGGAACC	TCTCAAGAAT	TCCCTCTTCAA	TACGACGAAA	CCATATATAA	ACAACATAAT	GATGGAATA	TTGATGAGCC		
TCTGTTGGCT	ACAATGATAT	ATATGTTTCA	TCGTGATCCA	CATCAAGTTT	TCCGTGAGCG	TATCGAACAG	GACGATGAGA		
ACTGGCAAAA	CATATTGCGC	ATATGTTTCA	TACGACGAAA	CCATATATAA	ACAACATAAT	GATGGAATA	TTGATGAGCC		
AAAGCAGTGA	ACACTTTGAA	ACAATTCAT	CATCGAATTG	GATGAACATG	CGATTCAAGC	CACCAACCAC	AGATGCTCCA		
GAAATCGGAT	GGAGAGTCGA	ATTCCGCGCA	ACTGAAGTTC	CAAGGCTGACA	TATTTGATGC	CAATTTCAT	GTTGCTTCGT		
TGTATTGCTC	ACCAGAATGA	TGATCTCCTT	AGAAATTCCT	GTTCAGAAAA	GGATTGGCTG	AGTGCAAATC	TGCTCCCGAA		
GTGCTCAGCA	AAAAGATGCA	GTTCTCAATC	ATGTGGACCA	CCTAGTCAAG	ATATTGAAGA	AATGTCGATT	GATGAGATTA		
AATTTGAAAG	GATCGGAGAA	ATGTGGACCA	CCTAGTCAAG	TATTCGCCAA	TTTCTAGATT	CTGCTGATGT	TGATGTGGAT		
GAAAAATGGA	TTCCCAGGTC	TCATTTCACT	TATTCGCCAA	TTTCTAGATT	TGGAGAGATT	AATACTTTGG	CTCACTGGAC		
CGATTTCTCA	ATATTGAAC	TTTATTTTGA	AACGAGCAAC	AGTGATGTAA	ATGATAATAT	AGTTTACGAT	CTTTTGAAAA		
GTACAATCTC	ATCCTGCATA	CAAACATGAC	AGTGATGTAA	ATGATAATAT	TGCTACCGCT	CTAAAACCGA	TCATGCCATT		
CATCTCAAAC	GGAGAAGATC	ACTGTGAGAA	GCTGCTCGGA	TGCTACCGCT	ACGTGCACAT	TAGgcgataa	ttgattgatt		
TTCGCAAAGC	TGAAGAGCAC	ATGATCGTGT	CCAGCCAAAA	ACGTGCACAT	TAGgcgataa	ttgattgatt	atgtgatttt		
aatttattta	tggtctatatac	gtcgtgtttc	ccattccttc	taggccttcc	atgattcaca	atgttctcat	gcatatcaaa		
tttagttggc	catctacatt	aaattactga	tatgtttgatg	ctatttttcta	gtaagcagat	gtcagtggtt	agtaattcaa		
aaatttaaac	tctgaatttc	taaatgcttg	ttttttgagt	agtaggaatc	gtacacgaatg	gtacattaat	ctgaaaataa		
tttcatattt	atgtacaatg	ctccccctgaa	tccatcatat	aattattatc	cgtgttg				

FIG. 28



## T19E7.2c (spliced)

```

aatcgttctt cttcttattt tctacagctg atgatgtttg atgaagggtt tatttttctt gctttttcca
ccctgttaat attattttcg atattcccaa aaataattcc aaattttcag tccatattca tctggatact
tgcaacatca tcaactgattt tggtagtcag ttcaccatcg tccaacacct caatccaatc atcgtcatac
gatcggatca cgacaaaaca tcttctggac aatatatcac cgacatttag tgagtatgac ttgaaaagtg
catctgatca cttttcgagc cgttttgtcg ctagggactt tttaatgaat cagatgtact ttctgaattt
tttagagcaa aagcagtagt tgcacttttg aaacttaaat taatatacaa aactatgata tatattttca
gaaatgtaca cggacagcaa taataggaac ttgatgaag tcaagcatca gcatcaacaa gaacaagatt
tcaatggcca atccaaatat gattatccac aattcaaccg tccaatgggt ctccgttggc gtgatgatca
acggatgatg gagtatttca tctcgaatcg tccagttagaa actgttccag ttatgccaat actcagcagc
catccaccag catctccatt cggtagagga ccattctacg aacgtccaac cagatcatct cgataccagt
acagttccgc ttctctcgag gatattcgact tgattgatgtg gctatggaga agtgatattg ctggagagaa
gggcagacga caagtggctc tctctgatca ctacgaatgt gatttccaga cgttgagaga gaaatcgaca
gtagccggcag tcaactggcga agagaatgct cgatatgaag atcttccga accattctat taatggattct
tccagtcctt caataacaa caatattcagc agaaagatga gaaacaaacaa ccagaacaaat taagacagcc
aactcttga caatccaaactc aaaaagccga atttgaagat gatctctttc atcaagatct tgcctcagctt
ttcaggagatg ttctacagaa agaagcagaa ttcaatcaac ttcttgataa taagcaacaa catccagtta
tcaataatgt ttctcttctc gaaggaattg ttatatatca cccaaatttc agcgagatgc aagagatccg
tgatttctgc aatcaagttt ccatttcaac aattccaaga acatccagctg ctcaaccaga gactttgttc
aatgtaacgg atttcacagac tgcgaacag tggcttccaa cagaagtgt accaaaccat gtgttcccaa
catccacta cccctacatt ccaatggcaa agcacactct tcaaccagct ctatcaaatg gagagattga
ctatgatcat tccatcaaat ccactcgtca gactccagct tctcgtctca tcaatccagct ttccaggagct
caacagcaga ctcaaacgag cccaggaagg ctacagctga ctgcaacagg tactcaatcg ttcttccagc
catatcactc acagagacac tccgtttagt pattggcagat tcaatccgtca tcaacgttct ctccgctctc
ttcggaaatct ccacgataga cgtcagagag ctcaaccgga actcaggagt ctcttttcta cggaaagtgc
gctccatcca gtggatcagg ctaccaagga tgatcgtctc caccgttcat acaatgttcc attaagattc
cgagagtgtt tccagtgccc agcggacaac cgaagcgttg aggtcaatcg aaggatgagc agctcccgag
tcacnaccag cttccagtgt cggcgttcca gatttgggag atgtcattaa gggagttgca acaagtcttg
aagaaccaga gtcctacgga gtatcaaga cagttgattc ccaagattcg tccagcggga aagacaagg
ttgcttcccg cacttggcgt caaagacgga cccatcgtca tgcagaagat tccgattaca tctgaagaagc
cctcttttat cacataaaat ctcggtcgaa acctatttaa agccacataa ttaaagataa ttaattccgc
cacaataate gtttttttct tctttgccgt gtctcatttc attttgatct actctttcct ccttcggat
tctttgattt cccagtgaaa tacctcacc ctttcaatcc ccacaaagtg agcaaccctt atcttgcaac
agttttatca tctcttcac ataccagtt tgataattta ttatctgat cccatccctt tgcgcctct
cattagatc ctagttttct atttgagccc ggagctcaga ctacatctcc gaatcatcat acaaatagat
agaaacgggt ctcgtagcga aagaatacgt gcaccacag acccccccac cctgttcacc cccataccc
tgaaaaatat gatctttaca gttatttcta ttatatctc aaatctctcg taatatcgta tcaattcct
cttctttttt gtcattttca atttttctca aattttctcg atctattctt tttcttgat ttttggaact
tgtatccctc ctccatcccc agact

```

## T19E7.2c (conceptual translation)

```

MYTDSNNRNF DEVNHQHQE QDFNGQSKYD YPQFNRPML RWRDDQRMME YFMSNGPVET VPVMPILTEH
PPASPFGRGP STERPTSSR YEYSSPSLED IDLIDVLWRS DIAGEKGTRQ VAPADQYECD LQTLTEKSTV
APLTAENAR YEDLSKGFYN GFFESFNNNQ YQKHQHQQR EQIKPTLEH PTQAELEDD LFDDELAQLF
EDVSREEGQL NQLFDNKQQH PVINNVSLSE GIVYNQANLT EMQEMRDSCN QVSISTIPTT STAQPETLFN
VTDSQTVEQW LPTEVPNDV FPTSNYAYIG MQNDLSLQAV SNGQIDYDHS YQSTGQTPLS PLIIGSSGRO
QQTQTSPPSV TVTATATQSL FDPYHSQRHS FSDCTTDSS TCSRLSSESP RYTSSESSTGT HESRFYGLA
PSSGSRYQRS SSPRSSQSSI KIARVVPLAS GQRKRGROSK DEQLASDNEL PVSAFQISEM SLSELQVVLK
NESLSEYQRQ LIRKIRRRGK NKVAARTCRQ RRTDRHDKMS HYI*

```

FIG. 29

## T19E7.2b (spliced)

```

gaATGTCACCT TCCATCTGAT TTGCTCTCT CTCTTCTGGC ATCCTCTACC ACCACCAACA CCACCAACAC
CGCTCCAGCA GCTGTCAACT CTTTIGACGA ACAAGAAGAA GAATCCAAGA AGATACTGAA CATGTACCTT
CAAATGTTCA ATCAACAACA GGTGGATCAA CACGGCCATC ATCACCACA TCCATACGCC TATTGAGGAG
TCTCGAGGAC TTTIGACAGA GTGTTCGCAATCATCCAACTATCGGCTAGATT CGAATGCCAAATACGACAGTCT
TCAAGCAGTT GTATCAATTC GACAGATTGAGCTATGATCAT TCGTATCATT CCAGTGGTCA GACTCCACTC
TGTGCTCTCA TCAATTCGATC TTGCTTCGATC CATATCAGTC AGAGAGACAG TCGTTTACGT ATTCGACTAG
CTGCAACAGC TACTCAATCG TTGTTTCGATC CATATCAGTC AGAGAGACAG TCGTTTACGT ATTCGACTAG
TGATTCTGCA TCAACGTGCT CTGCTCTCTC TTGCGGAATCT CCACGATACA CGTCAGAGAG CTCAACCGGA
ACTCAGGAGT CTGCTTCTA GGGAAAGTTG GCTCGATCCA GTGGATCACC CTACCAACGA TCATCGTCTC
GAGCTTCATC ACAATCTTCG ATTAAGATCG CGAGAGTGT TCCACTGGCC AGCGGAGAAC GGAAGCGTGG
ACGTCAATCG AAGGATGAGG AGCTCGCCAG TGACAACGAG CTTCAGTGT CCGCGTTCCA GATTTCGGAG
ATGTCATTAA GCGAGTTGCA ACAAGTGTG AAGAACGAGA GTCTCAGCGA GTATCAAGA CAGTTGATTG
GCAAGATTCT TCGACGCGCA AAGAACAAGG TTGCTGCGCG CAGTTGCGGT CAAGAGACCA CCGATCGTCA
CGACAAGATG TCGGATTAGA TCTGAgaagc cctcttttat cacataaaat ctcggtcgaa accttattaa
agccacataa ttaaagataa ttaattccgc cacaataatc gtttttttct tctttgccgt glctcatttc
atthtgatct actctttcct cccttcggat tctttgattt cccagtgaat tacctcacc acttcaatcc
ccacaagtg agcaacccct atcttgcaac agttttatca tctcttcac ataccagtt tgataattta
ttatctgatc cccatccct tgtcgctct ctagttttc atttgagccc ggagctcaga
ctacatctcc gaatcatcat acaaatagat agaaacgggt ctctgacga aagaatacgt gcaccacag
accccccat cctgttcacc cccatacacc tgaaaaatat gatctttaca gttatttcta ttatctctc
aatctctcg taatatcgta tcaatttct cttcttttt gtcattttca atttttctca aatttctcag
atctattctt tttcttgat ttttggaa tgtatccct cctccatccc agacttcccc ttcccagtta
ctcttgata ttttcatata tgtccatata tcgtttgaat ctctcattta

```

## T19E7.2b (conceptual translation)

```

MSLPDFASS LLASSTTTNT TNTAPAAVNS FDEQEEESKK ILNMYLQMFN QQQVDQHGHH HQHPYAYSGV
SSTFDRVFPT SNYAYIGMQN DSLQAVVNS QIDYDHSYQS TGQTPLSPLI IGSSGRQQQT QTSPPSVTVT
ATATQSLFDP YHSQRHSFSD CTTDSSSTCS RLSSSPRYT SESSTGTHES RFYGKLAPSS GSRYQRSSSP
RSSQSSIKIA RVVPLASQOR KRGRQSKDEQ LASDNELPVS AFQISEMSLS ELQOVLKNES LSEYQQLIR
KIRRRGKNKV AARTCRQRT DRHDKMSHYI *

```

FIG. 30

## T19E7.2a (spliced)

ATGGGCGGTT CATCAGCCG TCAGCGAAGT ACCTCGGCGA CGAGACGAGA CGATAAACGA AGACGAAGAC  
 AGTGCTTCTC TTCGGTAGCC GACGACGAAG AAGAGACGAC GTCAATTTAT GGAGTGTCCT GCATATTCAT  
 CTGGATACCT GCAACATCAT CACTGATTTT AGTGATCAGT TCACCATCGT CCAACACCTC AATCGAATCA  
 TCGTCATACG ATCGGATCAC GACAAACATC CTCTGGAGCA ATATATCAGG AGCATTTAAA ATGTACACGG  
 ACAGCAATAA TAGGAACCTT GATGAAGTCA ACCATCAGCA TCAACAAGAA CAAGATTICA ATGGCCAATC  
 GAAATATGAT TATCCACAAT TCACCCGTCC AATGGGTCTC CGTTGGGGTG ATGATGAACG GATGATGGAG  
 TATTTCATGT CGAATGGTCC AGTAGAAACT GTTCCAGTTA TGCCAATACT CACCGAGCAT CCACCAGCAT  
 CTCCATTCCG TAGAGGACGA TCTACAGAAC GTCCAAGGAC ATCATCTCGA TACGAGTACA GTTTCGCTTC  
 TCTCGAGGAT ATCGACTTGA TTGATCTGCT ATGGAGAAGT GATATTGCTC GAGAGAAGGG CACACGACAA  
 CTGGCTCCTG CTGATCAGTA CGAATGTGAT TTGCAACGCT TGACAGAGAA ATCGACAGTA GCGGCACTCA  
 CTGCGGAAGA GAATGCTCGA TATCAAGATC TTTCGAAAGG ATTCTATAAT CGATTCTTCG AGTGGTTCAA  
 TAACAATCAA TATCAGGAGA AACATCAGCA ACAACAAGCA GAAGAATAAA AGACAGCAAC TCTTGAACT  
 CGAAGTCAA AAAGCGAATT GCAACATGAT CTGTTTCAAT GCAACAAGAT CCAGTTATCA ATAATGTTTC  
 CAAGAGAAGA AGGAGCAATG AATCAACCTT TTGATATAAT GCAACAAGAT CCAGTTATCA ATAATGTTTC  
 TCTGTCGGAA CGAATCTTTC ATAATCAGCC AAATTCAGC GAGATCGAAG AGATGCGTGA TTCTGCAAT  
 CAAGTTTCCA TTTCACAAT TCCAACAACA TCGAGTGTCT AACGAGAGAC TTGTTTCAAT GTAACCGATT  
 CACAGACTGT CGAAGACTGG GTTCCAACAG AAGTGTACC AAAGCATCTG TTCCCAAGAT CCAACTAGCG  
 CTACATTCGA ATGCAAAACG ACAGCTTICA AGCAGTTGTA TCAATGCAAG AGATTGACTA TGATCATTCG  
 TATCAATCCA CTGGTCAGAC TCCACTCTCT CCTCTCATCA TTGGATCTTC AGGAGCTCAA CAGCAGCTG  
 AAAGGAGCCG AGGAGCCGTC ACAGTGAAGT CAAACAGTAC TCAATCGTTG TTGGATCCAT ATGACTCACA  
 GAGACACTCG TTACTGATT GCACTACTGA ATCTCATCAT AGGTGCTCTC GCCTCTCTTC GGAATCTCCA  
 CCATACAGCT CAGAGAGCTC AACCGGAAC CAGGACTCTC GTTCTACGG AAAGTTGGCT CCATCCAGTC  
 GATCAGCTA CCAACGATCA TCGTCTCCAC GTTCTACAGA ATCTTCGATT AAGATCGCGA GAGTTGTTCG  
 ACTGGCCAGC CGACAACGGA AGCGTGGACG TCAATCCAAG GATGAGCAGC TCGCCAGTGA CAACGAGCTT  
 CCAGTGTCCG CGTTCCAGAT TTGGGAGATG TCATTAAAGC AGTTGCAACA AGTGTGAAG AACGAGCTG  
 TCAGCGAGTA TCAAGACAGT TTGATTGGCA AGATTCTCG AGGCGAAG AACAAAGCTTC CTGCGCGAC  
 TTGCGCTCAA ACAGCCACCG ATCGTCAGCA CAGATCTCC CATTACATCT GAGAAGCCCT CTTTTATCAC  
 ataaaatctc ggtcgaaacc ttattaaagc cacataatta aagataatta attccgccac aataatcggt  
 tttttcttct ttgcogtgct tcatttcatt ttgatctact ctttctccc cttccgattct ttgatttccc  
 agtgaataac ctcacccact tcaatcccca caaagtgagc aaccctatc ttgcaacagt tttatcatct  
 cttcatcata cccagtttga taatttatta tctgatcccc atcccttgt cgctctcat tagtatccta  
 gtttttcatt tgagcccgga gtcagacta catctccgaa tcatcatata aatagataga aacgggtctc  
 gtgacgaaag aatacgtgca ccacacgacc ccccatcct gttcaccctc atacacctga aaaatatgat  
 ctttacagtt atttctatta tatctcaaa tctctcgtaa tatcgatca atttctctt ctttttggtc  
 attttcaatt tttctcaaat ttctcagatc tattcttttt cttgtatttt tggaaactgt atccctctc  
 catccccaga cttccctctc ccagttactc ttgtacattt tcatatatgt ccatatatcg tttgaatctc  
 tcattttatgg aaataaattt gaaaaaatc

## T19E7.2a (conceptual translation)

MGGSSRRQRS TSATRRDDKR RRRQCFSSVA DDEEETTSIY GVSSIFIWIL ATSSLILVIS SPSSNTSIQS  
 SSYDRITTKH LLDNISPTFK MYTDSNNRNF DEVNHQHQE QDFNGOSKYD YPQFNRPMLG RWRDDQRMME  
 YFMSNGPVET VPVMPILTEH PPASPFGRGP STERPTTSSR YEYSSPSLED IDLIDVLWRS DIAGEKGTRQ  
 VAPADQYECD LQTLTEKSTV APLTABENAR YEDLSKGFYN GPFESFNNNQ YQOKHQQQQR EQIKTPTLEH  
 PTQAELEDD LFDDELALQF EDVSREEGQL NQLFDNKQQH PVINNVLSE GIVYNQANLT EMQEMRDSCN  
 QVSISTIPTT STAQPETLFN VTDSQTVQW LPTEVVPNDV FPTSNYAYIG MONDSLQAVV SNGQIDYDHS  
 YQSTGQTPLS PLIIGSSGRQ QQTQTSPPSV TVTATATQSL FDPYHSQRHS FSDCTTDDSS TCSRLSSSEP  
 RYTSESTGT HESRFYGLA PSSGSRYQRS SSPRSSQSSI KIARVVPLAS GQRKRGRQSK DEQLASDNEI  
 PVSAFQISEM SLSELQVLK NESLSEYQRQ LIRKIRRRGK NKVAARTCRQ RRTDRHDKMS HYI\*

FIG. 31

# Document made available under the Patent Cooperation Treaty (PCT)

International application number: PCT/US04/019046

International filing date: 14 June 2004 (14.06.2004)

Document type: Certified copy of priority document

Document details: Country/Office: US  
Number: 60/478,185  
Filing date: 13 June 2003 (13.06.2003)

Date of receipt at the International Bureau: 23 March 2005 (23.03.2005)

Remark: Priority document submitted or transmitted to the International Bureau in compliance with Rule 17.1(a) or (b)



World Intellectual Property Organization (WIPO) - Geneva, Switzerland  
Organisation Mondiale de la Propriété Intellectuelle (OMPI) - Genève, Suisse

**This Page is Inserted by IFW Indexing and Scanning  
Operations and is not part of the Official Record**

**BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ **BLACK BORDERS**
- ☐ **IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- ☐ **FADED TEXT OR DRAWING**
- ☐ **BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- ☐ **SKEWED/SLANTED IMAGES**
- ☒ **COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- ☐ **GRAY SCALE DOCUMENTS**
- ☒ **LINES OR MARKS ON ORIGINAL DOCUMENT**
- ☐ **REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- ☐ **OTHER: \_\_\_\_\_**

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.**